

DEFINITION HS_5142_A1_D07_SP6E RPCI-11 Human Male BAC Library Homo sapiens
genomic clone Plate=718 Col=13 Row=G, DNA sequence.
ACCESSION AQ457751
VERSION AQ457751.1 GI:4636391
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 454)
AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T., and
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
TITLE Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
JOURNAL Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)
MEDLINE 99380589
COMMENT Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
or from Resear h Genetics (info@resgen.com). BAC end Web Server:
http://www.htsc.washington.edu
Plate: 718 row: G column: 13
Seq primer: SP6
Class: BAC ends
FEATURES
source
1..454
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=718 Col=13 Row=G"
/clone_lib="RPCI-11 Human Male BAC Library"
/sex="male"
/note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
Male blood DNA was isolated from one randomly chosen donor
and partially digested with a combination of EcoRI and
EcoRI Methylase. Size selected DNA was cloned into the
pBACe3.6 vector at EcoRI sites"
BASE COUNT 128 a 83 c 95 g 145 t 3 others
ORIGIN

Query Match 85.6%; Score 15.4; DB 97; Length 454;
Best Local Similarity 94.1%; Pred. No. 8.4e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 ctcccagcgtgcgccat 18
|||||
Db 300 CTCCCAGCGTGAGCCAT 316

Search completed: December 4, 2000, 21:06:39
Job time: 19239 sec

Email: Robert_Strausberg@nih.gov
This clone is available royalty-free through LLNL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Seq primer: -40UP from Gibco
High quality sequence stop: 456.

FEATURES
source

Location/Qualifiers
1. .525
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2662007"
/clone_lib="Soares_NFL_T_GBC_S1"
/lab_host="DH10B"
/note="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia) with
a modified polylinker; Site_1: Not I; Site_2: Eco RI;
Equal amounts of plasmid DNA from three normalized
libraries (fetal lung NbHL19W, testis NHT, and B-cell
NCI_CGAP_GCB1) were mixed, and ss circles were made in
vitro. Following HAP purification, this DNA was used as
tracer in a subtractive hybridization reaction. The driver
was PCR-amplified cDNAs from pools of 5,000 clones made
from the same 3 libraries. The pools consisted of
I.M.A.G.E. clones 297480-302087, 682632-687239,
726408-728711, and 729096-731399. Subtraction by Bento
Soares and M. Fatima Bonaldo. "
154 a 104 c 111 g 155 t 1 others

BASE COUNT
ORIGIN

Query Match 88.9%; Score 16; DB 20; Length 525;
Best Local Similarity 100.0%; Pred. No. 4.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 tcccgagcgtgcgccat 18
|||||
Db 305 TCCGAGCGTGCGCCAT 320

RESULT 13
AI982970

LOCUS AI982970 735 bp mRNA EST 31-AUG-1999
DEFINITION wt46b05.x1 NCI_CGAP_Pan1 Homo sapiens cDNA clone IMAGE:2510481 3'
similar to TR:O43326 O43326 HYPOTHETICAL 65.4 KD PROTEIN. ;, mRNA
sequence.

ACCESSION AI982970
VERSION AI982970.1 GI:5810189
KEYWORDS EST.
SOURCE human.

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 735)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550

JOURNAL
COMMENT

Email: Robert_Strausberg@nih.gov
Life Technologies catalog #: 11548-013
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Seq primer: -40UP from Gibco
High quality sequence stop: 426.

FEATURES
source

Location/Qualifiers
1. .735
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2510481"
/clone_lib="NCI_CGAP_Pan1"
/tissue_type="adenocarcinoma"
/lab_host="DH10B"

/note="Organ: pancreas; Vector: pCMV-SPORT6; Site_1: SalI;
Site_2: NotI; Cloned unidirectionally. Primer: Oligo dT.
Average insert size 1.72 kb. Life Technologies catalog #:
11548-013"
215 a 150 c 153 g 217 t

BASE COUNT
ORIGIN

Query Match 88.9%; Score 16; DB 14; Length 735;
Best Local Similarity 100.0%; Pred. No. 4.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 tcccgagcgtgcgccat 18
|||||
Db 304 TCCGAGCGTGCGCCAT 319

RESULT 14

AQ214736 416 bp DNA GSS 18-SEP-1998
LOCUS HS_3250_B1_F05_MR CIT Approved Human Genomic Sperm Library D Homo
DEFINITION sapiens genomic clone Plate=3250 Col=9 Row=L, DNA sequence.
ACCESSION AQ214736
VERSION AQ214736.1 GI:3625937
KEYWORDS GSS.
SOURCE human.

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 416)

REFERENCE

AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.

TITLE

Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)
99380589

JOURNAL

MEDLINE

COMMENT

Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Sequence Tagged Connector
Plate: 3250 row: L column: 9
Class: BAC ends
High quality sequence stop: 416.

FEATURES

source

Location/Qualifiers
1. .416
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=3250 Col=9 Row=L"
/clone_lib="CIT Approved Human Genomic Sperm Library D"
/sex="male"
/note="Organ: sperm; Vector: pBelobAC11; BAC Clones in
E-Coli DH10B"
120 a 123 c 72 g 100 t 1 others

BASE COUNT
ORIGIN

Query Match 85.6%; Score 15.4; DB 90; Length 416;
Best Local Similarity 94.1%; Pred. No. 8.3e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 ctcccgagcgtgcgccat 18
|||||
Db 177 CTACCAGCGTGCGCCAT 193

RESULT 15

AQ457751

LOCUS

AQ457751 454 bp DNA GSS 23-APR-1999

```

BASE COUNT      121 a      85 c      95 g      137 t      1 others
ORIGIN

Query Match      88.9%; Score 16; DB 19; Length 439;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 tccacgctgcgccat 18
    |||
Db 306 TCCACGCGTGC GCCAT 321

RESULT 10
AI005163
LOCUS      AI005163      459 bp      mRNA      EST      27-AUG-1998
DEFINITION      oui3c07.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone
IMAGE:1626156 3' similar to TR:Q99499 Q99499 DYNEIN-RELATED
PROTEIN. ;, mRNA sequence.
ACCESSION      AI005163
VERSION      AI005163.1 GI:3214673
KEYWORDS      EST.
SOURCE      human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE      1 (bases 1 to 459)
AUTHORS      NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE      National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
JOURNAL      Tumor Gene Index
COMMENT      Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert_Strausberg@nih.gov
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 703 Std Error: 0.00
Seq primer: primer name ambiguous
High quality sequence stop: 444.
FEATURES      Location/Qualifiers
1..459
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1626156"
/clone_lib="Soares_NFL_T_GBC_S1"
/lab_host="DH10B"
/note="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia) with
a modified polylinker; Site_1: Not I; Site_2: Eco RI;
Equal amounts of plasmid DNA from three normalized
libraries (fetal lung NbHL19W, testis NHT, and B-cell
NCI-CGAP_GCB1) were mixed, and ss circles were made in
vitro. Following HAP purification, this DNA was used as
tracer in a subtractive hybridization reaction. The driver
was PCR-amplified cDNAs from pools of 5,000 clones made
from the same 3 libraries. The pools consisted of
I.M.A.G.E. clones 297480-302087, 682632-687239,
726408-728711, and 729096-731399. Subtraction by Bento
Soares and M. Fatima Bonaldo."
BASE COUNT      128 a      83 c      96 g      152 t
ORIGIN

Query Match      88.9%; Score 16; DB 7; Length 459;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 tccacgctgcgccat 18
    |||
Db 350 TCCACGCGTGC GCCAT 365

RESULT 11
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AI682599      510 bp      mRNA      EST      17-DEC-1999
LOCUS      wc63ell.x1 NCI_CGAP_Pan1 Homo sapiens cDNA clone IMAGE:2323340 3'
DEFINITION      similar to TR:O43326 O43326 HYPOTHETICAL 65.4 KD PROTEIN. ;, mRNA
sequence.
ACCESSION      AI682599
VERSION      AI682599.1 GI:4892781
KEYWORDS      EST.
SOURCE      human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE      1 (bases 1 to 510)
AUTHORS      NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE      National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
JOURNAL      Tumor Gene Index
COMMENT      Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert_Strausberg@nih.gov
Life Technologies catalog #: 11548-013
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Insert Length: 1483 Std Error: 0.00
Seq primer: -40UP from Gibco
High quality sequence stop: 360.
FEATURES      Location/Qualifiers
1..510
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2323340"
/clone_lib="NCI_CGAP_Pan1"
/tissue_type="adenocarcinoma"
/lab_host="DH10B"
/note="Organ: pancreas; Vector: pCMV-SPORT6; Site_1: SalI;
Site_2: NotI; Cloned unidirectionally. Primer: Oligo dt.
Average insert size 1.72 kb. Life Technologies catalog #:
11548-013"
BASE COUNT      143 a      89 c      110 g      168 t
ORIGIN

Query Match      88.9%; Score 16; DB 12; Length 510;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 tccacgctgcgccat 18
    |||
Db 349 TCCACGCGTGC GCCAT 364

RESULT 12
AW182837
LOCUS      AW182837      525 bp      mRNA      EST      18-NOV-1999
DEFINITION      xj64dl2.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone
IMAGE:2662007 3' similar to TR:O43326 O43326 HYPOTHETICAL 65.4 KD
PROTEIN. ;, mRNA sequence.
ACCESSION      AW182837
VERSION      AW182837.1 GI:6451297
KEYWORDS      EST.
SOURCE      human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE      1 (bases 1 to 525)
AUTHORS      NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE      National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
JOURNAL      Tumor Gene Index
COMMENT      Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
```



```
RESULT 7
A1108632      658 bp      mRNA      EST      18-NOV-1998
LOCUS
DEFINITION   GH07838.5prime GH Drosophila melanogaster head pOT2 Drosophila
              melanogaster cDNA clone GH07838 5prime, mRNA sequence.
ACCESSION   A1108632
VERSION     A1108632.1  GI:3477167
SOURCE     EST.
           fruit fly.
ORGANISM   Drosophila melanogaster
           Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
           Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
           Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
REFERENCE   1 (bases 1 to 658)
AUTHORS    Harvey,D., Hong,L., Evans-Holm,M., Pendleton,J., Su,C., Brokstein
           ,P., Lewis,S. and Rubin,G.M.
TITLE      BDGP/HMI Drosophila EST Project
JOURNAL    Unpublished (1997)
COMMENT    Contact: Harvey, D.
           G. M. Rubin-Molecular and Cell Biology
           University of California Berkeley
           539 LSA, Berkeley, CA 94720-3200, USA
           Fax: 510 643 9947
           Email: http://www.fruitfly.org/EST, est@fruitfly.berkeley.edu
           Plate: 78 row: D column: 2
           High quality sequence stop: 520.
           Location/Qualifiers
             1..658
               /organism="Drosophila melanogaster"
               /db_xref="taxon:7227"
               /clone="GH07838"
               /sex="male and female"
               /dev_stage="adult"
               /lab_host="DH5 - alpha"
               /note="Organ: head; Vector: pOT2; Site_1: EcoRI; Site_2:
               XhoI; Sized fractionated cDNAs were directly ligated into
               pOT2. Plasmid cDNA library."
BASE COUNT  167 a  188 c  190 g  113 t
ORIGIN

Query Match      91.1%; Score 16.4; DB 8; Length 658;
Best Local Similarity 94.4%; Pred. No. 2.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
   |||
Db 80 TCGCCACGCGTGGCCAT 97

RESULT 8
AA990787      695 bp      mRNA      EST      24-NOV-1998
LOCUS
DEFINITION   LD34664.5prime LD Drosophila melanogaster embryo pOT2 Drosophila
              melanogaster cDNA clone LD34664 5prime, mRNA sequence.
ACCESSION   AA990787
VERSION     AA990787.1  GI:3177320
KEYWORDS    EST.
SOURCE     fruit fly.
ORGANISM   Drosophila melanogaster
           Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
           Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
           Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
REFERENCE   1 (bases 1 to 695)
AUTHORS    Harvey,D., Hong,L., Evans-Holm,M., Pendleton,J., Su,C., Brokstein
           ,P., Lewis,S. and Rubin,G.M.
TITLE      BDGP/HMI Drosophila EST Project
JOURNAL    Unpublished (1997)
COMMENT    Contact: Harvey, D.
           G. M. Rubin-Molecular and Cell Biology
           University of California Berkeley
```

```
539 LSA, Berkeley, CA 94720-3200, USA
Fax: 510 643 9947
Email: http://www.fruitfly.org/EST, est@fruitfly.berkeley.edu
Plate: 346 row: F column: 4
High quality sequence stop: 543.
Location/Qualifiers
  1..695
    /organism="Drosophila melanogaster"
    /db_xref="taxon:7227"
    /clone="LD34664"
    /clone_lib="LD Drosophila melanogaster embryo pOT2"
    /sex="male and female"
    /dev_stage="0 to 24 hours mixed stage embryonic"
    /lab_host="XLI Blue"
    /note="Organ: embryo; Vector: pOT2; Site_1: EcoRI; Site_2:
    XhoI; Sized fractionated cDNAs were directly ligated into
    pOT2."
BASE COUNT  178 a  201 c  206 g  110 t
ORIGIN

Query Match      91.1%; Score 16.4; DB 7; Length 695;
Best Local Similarity 94.4%; Pred. No. 2.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
   |||
Db 66 TCGCCACGCGTGGCCAT 83

RESULT 9
AW070294      439 bp      mRNA      EST      13-OCT-1999
LOCUS
DEFINITION   xa06g11.x1 Soares_NFL_T_GHC_S1 Homo sapiens cDNA clone
              IMAGE:2567588 3' similar to TR:O43326 O43326 HYPOTHETICAL 65.4 KD
              PROTEIN. ;, mRNA sequence.
ACCESSION   AW070294
VERSION     AW070294.1  GI:6025292
KEYWORDS    EST.
SOURCE     human.
ORGANISM   Homo sapiens
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (bases 1 to 439)
AUTHORS    NCI-CCGAP http://www.ncbi.nlm.nih.gov/ncicgap.
           National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
           Tumor Gene Index
JOURNAL    Unpublished (1997)
COMMENT    Contact: Robert Strausberg, Ph.D.
           Tel: (301) 496-1550
           Email: Robert_Strausberg@nih.gov
           This clone is available royalty-free through LLNL ; contact the
           IMAGE Consortium (info@image.llnl.gov) for further information.
           Seq primer: -40Up from Gibco
           High quality sequence stop: 400.
           Location/Qualifiers
             1..439
               /organism="Homo sapiens"
               /db_xref="taxon:9606"
               /clone="IMAGE:2567588"
               /clone_lib="Soares_NFL_T_GHC_S1"
               /lab_host="DH10B"
               /note="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia) with
               a modified polylinker; Site_1: Not I; Site_2: Eco RI;
               Equal amounts of plasmid DNA from three normalized
               libraries (fetal lung NbHL19W, testis NHT, and B-cell
               NCI_CGAP_GCE1) were mixed, and ss circles were made in
               vitro. Following HAP purification, this DNA was used as
               tracer in a subtractive hybridization reaction. The driver
               was PCR-amplified cDNAs from pools of 5,000 clones made
               from the same 3 libraries. The pools consisted of
               I.M.A.G.E. clones 297480-302087, 682632-687239,
               726408-728711, and 729096-731399. Subtraction by Bento
```

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810

Email: est@watson.wustl.edu
This clone is available royalty-free through LInL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 1759 Std Error: 0.00
Seq primer: mob.REGA+ET.

FEATURES

source

Location/Qualifiers
1. .442
/organism="Homo sapiens"
/db_xref="GDB:1278322"
/db_xref="taxon:9606"
/clone="IMAGE:361619"
/clone_lib="Soares retina N2b4HR"
/sex="male"
/tissue_type="retina"
/dev_stage="55 year old"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: eye; Vector: pT7T3D (Pharmacia) with a
modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st
strand cDNA was primed with a Not I - oligo(dT) primer [5'
TGTACCAATCTGAAGTGGAGCGCGCGCTTTTTTTTTT 3'],
double-stranded cDNA was size selected, ligated to Eco RI
adapters (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of a modified pT7T3 vector
(Pharmacia). The retinas were obtained from a 55 year old
Caucasian and total cellular poly(A)+ RNA was extracted 6
hrs after their removal. The retina RNA was kindly
provided by Roderick R. McInnes M.D. Ph.D. from the
University of Toronto. Library constructed by Bento
Soares and M.Fatima Bonaldo."

BASE COUNT 96 a 126 c 119 g 99 t 2 others
ORIGIN

Query Match 91.1%; Score 16.4; DB 40; Length 442;
Best Local Similarity 94.4%; Pred. No. 2.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 323 TCTCCACGCTGGGCCAT 306

RESULT 5

AI514786 577 bp mRNA EST 16-MAR-1999
LOCUS
DEFINITION LD46473.5prime LD Drosophila melanogaster embryo pOT2 Drosophila
melanogaster cDNA clone LD46473 5prime, mRNA sequence.
ACCESSION AI514786
VERSION AI514786.1 GI:4422958
KEYWORDS EST.
SOURCE fruit fly.
ORGANISM Drosophila melanogaster

Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

REFERENCE 1 (bases 1 to 577)
AUTHORS Harvey,D., Hong,L., Evans-Holm,M., Pendleton,J., Su,C., Brokstein

P., Lewis,S. and Rubin,G.M.
BDGP/HMI Drosophila EST Project
Unpublished (1997)
Contact: Harvey, D.

G. M. Rubin-Molecular and Cell Biology
University of California Berkeley
539 LSA, Berkeley, CA 94720-3200, USA
Fax: 510 643 9947

Email: http://www.fruitfly.org/EST, est@fruitfly.berkeley.edu
Plate: 464 row: G column: 1
High quality sequence stop: 458.

FEATURES

source

Location/Qualifiers
1. .577

/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/clone="LD46473"
/clone_lib="LD Drosophila melanogaster embryo pOT2"
/sex="male and female"
/dev_stage="0 to 24 hours mixed stage embryonic"
/lab_host="XL1 Blue"
/note="Organ: embryo; Vector: pOT2; Site_1: EcoRI; Site_2:
XhoI; Sized fractionated cDNAs were directly ligated into
pOT2."

BASE COUNT 140 a 175 c 170 g 92 t
ORIGIN

Query Match 91.1%; Score 16.4; DB 11; Length 577;
Best Local Similarity 94.4%; Pred. No. 2.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 73 TCGCCACGCTGGGCCAT 90

RESULT 6

AI512315 636 bp mRNA EST 16-MAR-1999
LOCUS
DEFINITION LD44124.5prime LD Drosophila melanogaster embryo pOT2 Drosophila
melanogaster cDNA clone LD44124 5prime, mRNA sequence.

ACCESSION AI512315
VERSION AI512315.1 GI:4421733
KEYWORDS EST.
SOURCE fruit fly.

ORGANISM

Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

REFERENCE

AUTHORS Harvey,D., Hong,L., Evans-Holm,M., Pendleton,J., Su,C., Brokstein
P., Lewis,S. and Rubin,G.M.
BDGP/HMI Drosophila EST Project
Unpublished (1997)
Contact: Harvey, D.

G. M. Rubin-Molecular and Cell Biology
University of California Berkeley
539 LSA, Berkeley, CA 94720-3200, USA
Fax: 510 643 9947

Email: http://www.fruitfly.org/EST, est@fruitfly.berkeley.edu
Plate: 441 row: B column: 12
High quality sequence stop: 557.

FEATURES

source

Location/Qualifiers
1. .636

/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/clone="LD44124"

/clone_lib="LD Drosophila melanogaster embryo pOT2"
/sex="male and female"
/dev_stage="0 to 24 hours mixed stage embryonic"
/lab_host="XL1 Blue"

/note="Organ: embryo; Vector: pOT2; Site_1: EcoRI; Site_2:
XhoI; Sized fractionated cDNAs were directly ligated into
pOT2."

BASE COUNT 158 a 188 c 184 g 106 t
ORIGIN

Query Match 91.1%; Score 16.4; DB 11; Length 636;
Best Local Similarity 94.4%; Pred. No. 2.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 77 TCGCCACGCTGGGCCAT 94

VERSION H83468.1 GI:1062139
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheraia; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 398)
Hillier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M., Holman
,M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J.,
Rifkin,L., Rohlfing,T., Soares,M., Tan,F., Trevaskis,E., Waterston
,R., Williamson,A., Wohldmann,P. and Wilson,R.
The WashU-Merck EST Project
Unpublished (1995)
JOURNAL Contact: Wilson RK
COMMENT Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
Source: IMAGE Consortium, LLNL
This clone is available royalty-free through LLNL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 1825 Std Error: 0.00
Seq primer: M13RP1
High quality sequence stop: 305.
FEATURES Location/Qualifiers
source 1..398
/organism="Homo sapiens"
/db_xref="GDB:3850898"
/db_xref="taxon:9606"
/clone="IMAGE:222137"
/clone_lib="Soares retina N2b5HR"
/sex="male"
/tissue_type="retina"
/dev_stage="55 year old"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: eye; Vector: pT7T3D (Pharmacia) with a
modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st
strand cDNA was primed with a Not I - oligo(dT) primer [5'
TGTTACCAATCTGAAGTGGAGCGCGCTTTTCTTTTCTTTTCTTTT 3'],
double-stranded cDNA was size selected, ligated to Eco RI
adapters (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of a modified pT7T3 vector
(Pharmacia). The retinas were obtained from a 55 year old
Caucasian and total cellular poly(A)+ RNA was extracted 6
hrs after their removal. The retina RNA was kindly
provided by Roderick R. McInnes M.D. Ph.D. from the
University of Toronto. Library constructed by Bento
Soares and M.Fatima Bonaldo."
BASE COUNT 85 a 104 c 108 g 98 t 3 others
ORIGIN
Query Match 91.1%; Score 16.4; DB 37; Length 398;
Best Local Similarity 94.4%; Pred. No. 2.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 tctcccagcgtgcgccat 18
|||||
Db 324 TCTCCAGCGTGGGCCAT 307
RESULT 3
AA402089
LOCUS AA402089 412 bp mRNA EST 16-MAY-1997
DEFINITION zu53f02.r1 Soares ovary tumor NbHOT Homo sapiens cDNA clone
IMAGE:741723 5' similar to SW:YK18_YEAST P36132 HYPOTHETICAL 46.6
KD PROTEIN IN DAL80-GAP1 INTERGENIC REGION. ; mRNA sequence.
ACCESSION AA402089
VERSION AA402089.1 GI:2056072
KEYWORDS EST.
SOURCE human.

ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheraia; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 412)
Hillier,L., Allen,M., Bowles,L., Dubuque,T., Geisel,G., Jost,S.,
Kucaba,T., Lacy,M., Le,N., Lennon,G., Marra,M., Martin,J., Moore,B.,
Schellenberg,K., Steptoe,M., Tan,F., Theising,B., White,Y., Wylie
,T., Waterston,R. and Wilson,R.
WashU-Merck EST Project 1997
Unpublished (1997)
JOURNAL Contact: Wilson RK
COMMENT Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
This clone is available royalty-free through LLNL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Seq primer: -28m13 rev2 ET from Amer sham
High quality sequence stop: 378.
FEATURES Location/Qualifiers
source 1..412
/organism="Homo sapiens"
/db_xref="GDB:5941912"
/db_xref="taxon:9606"
/clone="IMAGE:741723"
/clone_lib="Soares ovary tumor NbHOT"
/sex="Female"
/tissue_type="ovarian tumor"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: ovary; Vector: pT7T3D (Pharmacia) with a
modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st
strand cDNA was primed with a Not I - oligo(dT) primer [5'
TGTTACCAATCTGAAGTGGAGCGCGCTTTTCTTTTCTTTTCTTTT 3'],
double-stranded cDNA was size selected, ligated to Eco RI
adapters (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of a modified pT7T3 vector
(Pharmacia). Library constructed by Bento Soares and
M.Fatima Bonaldo."
BASE COUNT 77 a 115 c 131 g 89 t
ORIGIN
Query Match 91.1%; Score 16.4; DB 4; Length 412;
Best Local Similarity 94.4%; Pred. No. 2.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 tctcccagcgtgcgccat 18
|||||
Db 77 TCTCCAGCGTCCGCCAT 94
RESULT 4
W96258/c
LOCUS W96258 442 bp mRNA EST 29-NOV-1996
DEFINITION ze42b06.r1 Soares retina N2b4HR Homo sapiens cDNA clone
IMAGE:361619 5', mRNA sequence.
ACCESSION W96258
VERSION W96258.1 GI:1426165
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheraia; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 442)
Hillier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M., Holman
,M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J.,
Rifkin,L., Rohlfing,T., Soares,M., Tan,F., Trevaskis,E., Waterston
,R., Williamson,A., Wohldmann,P. and Wilson,R.
The WashU-Merck EST Project
Unpublished (1995)
JOURNAL Contact: Wilson RK
COMMENT Washington University School of Medicine

117: em_gss6:*
118: em_gss7:*
119: em_gss8:*
120: em_gss9:*
121: em_gss10:*
122: em_gss11:*
123: em_gss12:*
124: em_gss13:*
125: em_gss14:*
126: em_gss15:*
127: em_gss16:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB	ID	Description
C 1	16.4	91.1	387	37	H85916	H85916 YS95C07.r1
C 2	16.4	91.1	398	37	H83468	H83468 YS91B09.r1
C 3	16.4	91.1	412	4	AA402089	AA402089 zu53f02.r
C 4	16.4	91.1	442	40	W96258	W96258 ze42b06.r1
5	16.4	91.1	577	11	AI514786	AI514786 LD46473.5
6	16.4	91.1	636	11	AI512315	AI512315 LD44124.5
7	16.4	91.1	658	8	AI108632	AI108632 GH07838.5
8	16.4	91.1	695	7	AA990787	AA990787 LD34664.5
9	16	88.9	439	19	AW070294	AW070294 xa06g11.x
10	16	88.9	459	7	AI005163	AI005163 oul3c07.x
11	16	88.9	510	12	AI682599	AI682599 wc63e11.x
12	16	88.9	525	20	AW182837	AW182837 xj64d12.x
13	16	88.9	735	14	AI982970	AI982970 wt46b05.x
14	15.4	85.6	416	90	AQ214736	AQ214736 HS_3250_B
15	15.4	85.6	454	97	AQ457751	AQ457751 HS_5142_A
16	15.4	85.6	467	100	AQ649394	AQ649394 Sheared D
17	15.4	85.6	601	115	FR0021601	AL014474 F.rubripe
18	15.4	85.6	686	100	AQ655156	AQ655156 Sheared D
19	15.4	85.6	783	113	CNS01T6S	AL166141 Tetraodon
20	15.4	85.6	1027	114	CNS04CRA	AL284815 Tetraodon
21	15	83.3	231	5	AA625067	AA625067 af66g07.r
22	15	83.3	719	114	CNS049ZN	AL281228 Tetraodon
23	15	83.3	871	36	BE545551	BE545551 601070216
24	14.8	82.2	168	108	AZ208425	AZ208425 SP_0150_A
25	14.8	82.2	185	4	AA421434	AA421434 zu27g02.r
26	14.8	82.2	191	103	AQ843749	AQ843749 LMAJFV1_1
27	14.8	82.2	224	6	AA853590	AA853590 NHTCae06
28	14.8	82.2	246	20	AW183217	AW183217 xj74a08.x
29	14.8	82.2	254	16	AV134225	AV134225 AV134225
30	14.8	82.2	310	104	AQ911626	AQ911626 LMAJFV1_1
31	14.8	82.2	346	4	AA465312	AA465312 aa24b11.r
32	14.8	82.2	363	106	AZ043716	AZ043716 RPCI-23-3
33	14.8	82.2	380	103	AQ843748	AQ843748 LMAJFV1_1
34	14.8	82.2	413	32	BE008986	BE008986 CM4-BN016
35	14.8	82.2	413	33	BE119385	BE119385 UI-R-CA0-
36	14.8	82.2	426	3	AA364148	AA364148 EST74711
37	14.8	82.2	428	14	AL040548	AL040548 DKFZp434I
38	14.8	82.2	437	115	FR0008579	Z92389 F.rubripes
39	14.8	82.2	442	21	AW246200	AW246200 2821849.5
40	14.8	82.2	443	34	BE255160	BE255160 601113191
41	14.8	82.2	467	34	BE264833	BE264833 601194308
42	14.8	82.2	475	38	N28555	N28555 yx37f03.r1
43	14.8	82.2	487	5	AA605873	AA605873 fa20g10.s
44	14.8	82.2	511	24	AW719286	AW719286 LjNEST_C1
45	14.8	82.2	522	98	AQ502968	AQ502968 V47D5 mtm

ALIGNMENTS

RESULT 1
H85916/c

LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

H85916 387 bp mRNA EST 21-NOV-1995
ys95c07.r1 Soares retina N2b5HR Homo sapiens cDNA clone
IMAGE:222540 5', mRNA sequence.
H85916
H85916.1 GI:1067495
EST.
human.
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 387)
Hillier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M., Holman
,M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J.,
Rifkin,L., Rohlfig,T., Soares,M., Tan,F., Trevaskis,E., Waterston
,R., Williamson,A., Wohldmann,P. and Wilson,R.
The WashU-Merck EST Project
Unpublished (1995)
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: estewatson.wustl.edu
High quality sequence stops: 305
Source: IMAGE Consortium, LLNL
This clone is available royalty-free through LLNL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 1370 Std Error: 0.00
Seq primer: M13RP1
High quality sequence stop: 305.

FEATURES
source
Location/Qualifiers
1. .387
/organism="Homo sapiens"
/db_xref="GDB:3851301"
/db_xref="taxon:9606"
/clone="IMAGE:222540"
/clone_lib="Soares retina N2b5HR"
/sex="male"
/tissue_type="retina"
/dev_stage="55 year old"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: eye; Vector: pT7T3D (Pharmacia) with a
modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st
strand cDNA was primed with a Not I - oligo(dT) primer [5'
TGTTACCAATCTGAAGTGGGAGCGCGCGCTTTTTTT 3'] ,
double-stranded cDNA was size selected, ligated to Eco RI
adapters (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of a modified pT7T3 vector
(Pharmacia). The retinas were obtained from a 55 year old
Caucasian and total cellular poly(A)+ RNA was extracted 6
hrs after their removal. The retina RNA was kindly
provided by Roderick R. McInnes M.D. Ph.D. from the
University of Toronto. Library constructed by Bento
Soares and M.Fatima Bonaldo."

BASE COUNT
ORIGIN

82 a 100 c 107 g 95 t 3 others

Query Match 91.1%; Score 16.4; DB 37; Length 387;
Best Local Similarity 94.4%; Pred. No. 2.6e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 327 TCTCCACGCTGGCCCAT 310

RESULT 2
H83468/c

LOCUS
DEFINITION
ACCESSION
H83468 398 bp mRNA EST 13-NOV-1995
ys91b09.r1 Soares retina N2b5HR Homo sapiens cDNA clone
IMAGE:222137 5', mRNA sequence.
H83468

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: December 4, 2000, 15:46:00 ; Search time 1141.84 Seconds
(without alignments)
97.466 Million cell updates/sec

Title: US-09-369-941-1
Perfect score: 18
Sequence: 1 tctcccagcgtgcgccat 18

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 7189864 seqs, 3091403243 residues

Total number of hits satisfying chosen parameters: 14379728

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : EST:*
1: gb_est1:*
2: gb_est2:*
3: gb_est3:*
4: gb_est4:*
5: gb_est5:*
6: gb_est6:*
7: gb_est7:*
8: gb_est8:*
9: gb_est9:*
10: gb_est10:*
11: gb_est11:*
12: gb_est12:*
13: gb_est13:*
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42: em_estfun:*
43: em_esthum1:*

44: em_esthum2:*
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66: em_estin4:*
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86: em_estrol2:*
87: em_estrol3:*
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91: gb_gss4:*
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93: em_gss2:*
94: em_gss3:*
95: em_gss4:*
96: gb_gss5:*
97: gb_gss6:*
98: gb_gss7:*
99: gb_gss8:*
100: gb_gss9:*
101: gb_gss10:*
102: gb_gss11:*
103: gb_gss12:*
104: gb_gss13:*
105: gb_gss14:*
106: gb_gss15:*
107: gb_gss16:*
108: gb_gss17:*
109: gb_gss18:*
110: gb_gss19:*
111: gb_gss20:*
112: gb_gss21:*
113: gb_gss22:*
114: gb_gss23:*
115: gb_gss24:*
116: em_gss5:*

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 4, 2000, 15:46:00 ; Search time 993.06 Seconds
(without alignments)
79.166 Million cell updates/sec

Title: us-09-369-941-l
Perfect score: 18
Sequence: 1 tctcccagcgtgcgccat 18

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1033670 seqs, 2183789903 residues 2067340

Total number of hits satisfying chosen parameters:

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : GenEmbl:
1: gb_ba1:*
2: gb_ba2:*
3: gb_om:*
4: gb_ov:*
5: gb_pat:*
6: gb_ph:*
7: gb_pl1:*
8: gb_pl2:*
9: gb_pr1:*
10: gb_pr2:*
11: gb_pr3:*
12: gb_ro:*
13: gb_sy:*
14: gb_un:*
15: em_fun:*
16: em_hum1:*
17: em_hum2:*
18: em_in:*
19: em_om:*
20: em_or:*
21: em_ov:*
22: em_pat:*
23: em_ph:*
24: em_pl:*
25: em_ro:*
26: em_sts:*
27: em_sy:*
28: em_un:*
29: em_vi:*
30: gb_ba3:*
31: gb_in1:*
32: gb_in2:*
33: gb_in3:*
34: gb_pl3:*
35: gb_pr4:*
36: em_ba1:*
37: em_ba2:*
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39: em_htg2:*
40: em_htg3:*
41: em_htg4:*
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em_htg7:*
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62: em_hum5:*
63: em_hum6:*
64: gb_pr5:*
65: gb_pr6:*
66: gb_pr7:*
67: gb_htg1:*
68: gb_htg2:*
69: gb_htg3:*
70: gb_htg4:*
71: gb_htg5:*
72: gb_htg6:*
73: gb_htg7:*
74: gb_htg8:*
75: gb_htg9:*
76: gb_htg10:*
77: gb_htg11:*
78: gb_htg12:*
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85: gb_htg19:*
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87: gb_htg21:*
88: gb_htg22:*
89: gb_htg23:*
90: gb_sts1:*
91: gb_sts2:*
92: gb_vil:*
93: gb_vi2:*
94:

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	18	100.0	18	5	AR052619 Sequence
2	18	100.0	18	5	AR052624 Sequence
3	18	100.0	18	5	I96098 Sequence 17
c 4	18	100.0	27	5	AR004426 Sequence
c 5	18	100.0	27	5	I43661 Sequence 13
c 6	18	100.0	27	5	I86720 Sequence 8
c 7	18	100.0	35	5	AR052604 Sequence
c 8	18	100.0	35	5	I96083 Sequence 2
c 9	18	100.0	454	67	S72602 bcl2 [human
c 10	18	100.0	615	5	AR052623 Sequence
c 11	18	100.0	717	5	AR052622 Sequence
c 12	18	100.0	760	5	AR021160 Sequence

C 13 18 100.0 765 5 A76121
C 14 18 100.0 911 67 HUMBCL2B
C 15 18 100.0 1846 5 AR054009
C 16 18 100.0 1846 66 HSBCL2IG
C 17 18 100.0 5086 5 AR052621
C 18 18 100.0 5086 5 AR054008
C 19 18 100.0 5086 67 HUMBCL2A
C 20 18 100.0 5105 5 I08038
C 21 18 100.0 6030 67 HUMBCL2C
C 22 18 100.0 86692 75 AC021803
C 23 18 100.0 169542 89 AP001915
C 24 18 100.0 188601 69 AC009267
C 25 17 94.4 17 5 I96090
C 26 17 94.4 35737 9 AC005263
C 27 16.4 91.1 67568 74 AC020378
C 28 16.4 91.1 99614 68 AC008358
C 29 16.4 91.1 128132 68 AC007732
C 30 16.4 91.1 161799 9 AC002091
C 31 16.4 91.1 170757 68 AC007608
C 32 16.4 91.1 186323 68 AC006491
C 33 16.4 91.1 210299 31 AE003690
C 34 16.4 91.1 217026 72 AC013669
C 35 16 88.9 157432 83 AC069438
C 36 16 88.9 158934 81 AC055780
C 37 16 88.9 211341 85 AL138898
C 38 15.4 85.6 10854 1 AE001886
C 39 15.4 85.6 38616 9 AC004208
C 40 15.4 85.6 42245 9 AC004476
C 41 15.4 85.6 75163 83 AC069103
C 42 15.4 85.6 75533 68 AC006412
C 43 15.4 85.6 112630 78 AC025461
C 44 15.4 85.6 117143 67 HUMBRCA1
C 45 15.4 85.6 120841 10 AC008162

VERSION AR052624.1 GI:5975988
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 18)
AUTHORS Reed,J.C.
TITLE Regulation of bcl-2 gene expression
JOURNAL Patent: US 5831066-A 24 03-NOV-1998;
FEATURES Location/Qualifiers
source 1..18
BASE COUNT 2 a 8 c 4 g 4 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 18;
Best Local Similarity 100.0%; Pred. No. 83;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccaat 18
|||||
Db 1 TCTCCCAGCGTGCGCCAT 18

RESULT 3
I96098
LOCUS I96098 18 bp DNA
DEFINITION Sequence 17 from patent US 5734033.
ACCESSION I96098
VERSION I96098.1 GI:3940568
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.

PAT 01-DEC-1998

ALIGNMENTS

RESULT 1
AR052619
LOCUS AR052619 18 bp DNA
DEFINITION Sequence 17 from patent US 5831066.
ACCESSION AR052619
VERSION AR052619.1 GI:5975983
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE Unclassified.
AUTHORS Reed,J.C.
TITLE Regulation of bcl-2 gene expression
JOURNAL Patent: US 5831066-A 17 03-NOV-1998;
FEATURES Location/Qualifiers
source 1..18
BASE COUNT 2 a 8 c 4 g 4 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 18;
Best Local Similarity 100.0%; Pred. No. 83;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccaat 18
|||||
Db 1 TCTCCCAGCGTGCGCCAT 18

RESULT 4
AR004426/c
LOCUS AR004426 27 bp DNA
DEFINITION Sequence 13 from patent US 5747245.
ACCESSION AR004426
VERSION AR004426.1 GI:3965305
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.

PAT 04-DEC-1998

Query Match 100.0%; Score 18; DB 5; Length 18;
Best Local Similarity 100.0%; Pred. No. 83;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccaat 18
|||||
Db 1 TCTCCCAGCGTGCGCCAT 18

RESULT 2
AR052624
LOCUS AR052624 18 bp DNA
DEFINITION Sequence 24 from patent US 5831066.
ACCESSION AR052624
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE Unclassified.
AUTHORS Reed,J.C. and Sato,T.
TITLE Nucleic acids encoding Fas associated proteins and screening assays
JOURNAL Patent: US 5747245-A 13 05-MAY-1998;
FEATURES Location/Qualifiers
source 1..27
/organism="unknown"

BASE COUNT 7 a 6 c 10 g 4 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 27;
Best Local Similarity 100.0%; Pred. No. 77;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 25 TCTCCCAGCGTGCGCCAT 8

RESULT 5
I43661/c 27 bp DNA PAT 07-OCT-1997
LOCUS I43661 Sequence 13 from patent US 5632994.
DEFINITION I43661
ACCESSION I43661
VERSION I43661.1 GI:2468759
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 27)
AUTHORS Reed,J.C. and Sato,T.
TITLE Fas associated proteins
JOURNAL Patent: US 5632994-A 13 27-MAY-1997;
FEATURES Location/Qualifiers
source 1..27
/organism="unknown"

BASE COUNT 7 a 6 c 10 g 4 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 27;
Best Local Similarity 100.0%; Pred. No. 77;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 25 TCTCCCAGCGTGCGCCAT 8

RESULT 6
I86720/c 27 bp DNA PAT 10-JUN-1998
LOCUS I86720 Sequence 8 from patent US 5702897.
DEFINITION I86720
ACCESSION I86720
VERSION I86720.1 GI:3206438
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 27)
AUTHORS Reed,J.C. and Sato,T.
TITLE Interaction of proteins involved in a cell death pathway
JOURNAL Patent: US 5702897-A 8 30-DEC-1997;
FEATURES Location/Qualifiers
source 1..27
/organism="unknown"

BASE COUNT 7 a 6 c 10 g 4 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 27;
Best Local Similarity 100.0%; Pred. No. 77;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 25 TCTCCCAGCGTGCGCCAT 8

RESULT 7
AR052604/c 35 bp DNA PAT 29-SEP-1999
LOCUS AR052604 Sequence 2 from patent US 5831066.
DEFINITION AR052604
ACCESSION AR052604
VERSION AR052604.1 GI:5975968
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 35)
AUTHORS Reed,J.C.
TITLE Regulation of bcl-2 gene expression
JOURNAL Patent: US 5831066-A 2 03-NOV-1998;
FEATURES Location/Qualifiers
source 1..35
/organism="unknown"

BASE COUNT 6 a 8 c 13 g 8 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 35;
Best Local Similarity 100.0%; Pred. No. 73;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 35 TCTCCCAGCGTGCGCCAT 18

RESULT 8
I96083/c 35 bp DNA PAT 01-DEC-1998
LOCUS I96083 Sequence 2 from patent US 5734033.
DEFINITION I96083
ACCESSION I96083
VERSION I96083.1 GI:3940553
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 35)
AUTHORS Reed,J.
TITLE Antisense oligonucleotides inhibiting human bcl-2 gene expression
JOURNAL Patent: US 5734033-A 2 31-MAR-1998;
FEATURES Location/Qualifiers
source 1..35
/organism="unknown"

BASE COUNT 6 a 8 c 13 g 8 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 35;
Best Local Similarity 100.0%; Pred. No. 73;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 35 TCTCCCAGCGTGCGCCAT 18

RESULT 9
S72602/c 454 bp DNA PRI 07-MAY-1993
LOCUS S72602 bcl2 [human, 697 pre-B cell acute lymphocytic leukemia cell line,
DEFINITION Genomic, 454 nt].
ACCESSION S72602
VERSION S72602.1 GI:241046
KEYWORDS
SOURCE human 697 pre-B cell acute lymphocytic leukemia cell line.
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 454)

AUTHORS Tanaka,S., Louie,D.C., Kant,J.A. and Reed,J.C.
TITLE Frequent incidence of somatic mutations in translocated BCL2 oncogenes of non-Hodgkin's lymphomas
JOURNAL Blood 79 (1), 229-237 (1992)
MEDLINE 92096610
REMARK GenBank staff at the National Library of Medicine created this entry [NCBI gibbsq 72602] from the original journal article.

FEATURES
source 1. .454
/organism="Homo sapiens"
/db_xref="taxon:9606"
gene 1. .454
/partial
/gene="bcl2"
CDS 41. .433
/partial
/gene="bcl2"
/note="Unknown"
/codon_start=1
/protein_id="AAD14111.1"
/db_xref="GI:4261811"
/translation="MAHAGRTGYDNR EIVMKYIHYKLSQRGYENDAGDVGAAPPGAAP
APGIFSSQPGHTPHPAASRDVPARTSPLOTPAAPGAAGPALSPVPPVVHLALRQAGD
DFSRRYRGDFAEMSSQLHLTPFTARGRFA"

BASE COUNT 65 a 170 c 150 g 69 t
ORIGIN

Query Match 100.0%; Score 18; DB 67; Length 454;
Best Local Similarity 100.0%; Pred. No. 46;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 58 TCTCCCAGCGTGCGCCAT 41

RESULT 10
AR052623/c
LOCUS AR052623 615 bp DNA
DEFINITION Sequence 22 from patent US 5831066.
ACCESSION AR052623
VERSION AR052623.1 GI:5975987
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 615)
AUTHORS Reed,J.C.
TITLE Regulation of bcl-2 gene expression
JOURNAL Patent: US 5831066-A 22 03-NOV-1998;
FEATURES Location/Qualifiers
source 1. .615
/organism="unknown"
BASE COUNT 98 a 203 c 213 g 101 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 615;
Best Local Similarity 100.0%; Pred. No. 43;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 18 TCTCCCAGCGTGCGCCAT 1

RESULT 11
AR052622/c
LOCUS AR052622 717 bp DNA
DEFINITION Sequence 20 from patent US 5831066.
ACCESSION AR052622
VERSION AR052622.1 GI:5975986

KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 717)
AUTHORS Reed,J.C.
TITLE Regulation of bcl-2 gene expression
JOURNAL Patent: US 5831066-A 20 03-NOV-1998;
FEATURES Location/Qualifiers
source 1. .717
/organism="unknown"
BASE COUNT 113 a 237 c 237 g 130 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 717;
Best Local Similarity 100.0%; Pred. No. 42;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 18 TCTCCCAGCGTGCGCCAT 1

RESULT 12
AR021160/c
LOCUS AR021160 760 bp DNA
DEFINITION Sequence 11 from patent US 5789389.
ACCESSION AR021160
VERSION AR021160.1 GI:3975775
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 760)
AUTHORS Tarasiewicz,D.G., Schott,B., Holzmayer,T.A. and Roninson,I.B.
TITLE BCL2 derived genetic elements associated with sensitivity to chemotherapeutic drugs
JOURNAL Patent: US 5789389-A 11 04-AUG-1998;
FEATURES Location/Qualifiers
source 1. .760
/organism="unknown"
BASE COUNT 122 a 250 c 246 g 142 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 760;
Best Local Similarity 100.0%; Pred. No. 42;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 38 TCTCCCAGCGTGCGCCAT 21

RESULT 13
A76121/c
LOCUS A76121 765 bp DNA
DEFINITION Sequence 1 from Patent WO9320200.
ACCESSION A76121
VERSION A76121.1 GI:6088257
KEYWORDS human.
SOURCE Homo sapiens
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 765)
AUTHORS Evan,G.I.
TITLE MODIFIED CELLS AND METHOD OF TREATMENT
JOURNAL Patent: WO 9320200-A 14-OCT-1993;
IMP CANCER RES TECH (GB); EVAN GERARD IAN (GB)
FEATURES Location/Qualifiers
source 1. .765

/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="18"
31..750
/note="unnamed protein product"
/codon_start=1
/protein_id="CAB58588.1"
/db_xref="GI:6088258"
/translation="MAHAGRTGYDNRREIVMKYIHYKLSQRYEWDAGDVGAAPPGAAP
APGIFSSQPGHTPHPAASRDPVARTPLQTPAAPGAAAGPALSVPVPVHLLALRQAGD
DFSRRYRGDFAEMSSQLHLTPFTARGFATVVEELFRDGVNWGRIVAFFEFGVMCVE
SVNREMSPLVDNIALWMTEYLNRLHHTWIQDNGGWDAFVELYGPSMRPLFDDFSWSLSK
TLLSLALVGACITLGAYLSHK"
BASE COUNT 120 a 251 c 250 g 144 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 765;
Best Local Similarity 100.0%; Pred. No. 42;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 48 TCTCCAGCGTGCGCCAT 31

RESULT 14
HUMBCL2B/c
LOCUS HUMBCL2B 911 bp mRNA PRI 31-OCT-1994
DEFINITION Human B-cell leukemia/lymphoma 2 (bcl-2) proto-oncogene mRNA
encoding bcl-2-beta protein, complete cds.
ACCESSION M13995
VERSION M13995.1 GI:179368
KEYWORDS alternative splicing; bcl-2-beta protein; proto-oncogene.
SOURCE Human pre-B-cell leukemia cell line 380, cDNA to mRNA, clones B[15,16]; and DNA, clone lambda-18-27.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 911)
Tsujimoto,Y. and Croce,C.M.
Analysis of the structure, transcripts, and protein products of
bcl-2, the gene involved in human follicular lymphoma
Proc. Natl. Acad. Sci. U.S.A. 83 (14), 5214-5218 (1986)
86259760
Clean copy sequence for [1] kindly provided by Y.Tsujimoto,
10-FEB-1987. The bcl-2 gene is transcribed by alternative splicing
into three mRNAs of different sizes. It consists of at least two
exons and encodes two proteins which only differ at their
carboxy-terminal ends, and it is activated by translocation into
proximity with the Ig heavy chain locus. Both the normal and
rearranged bcl-2 gene products are expressed in the B-cell
leukemia/lymphoma 2 cells. Genomic clone lambda-18-27 contained
all the DNA sequences on the 5' of the splice site (position 732).
Location/Qualifiers
1..911
/organism="Homo sapiens"
/db_xref="taxon:9606"
/map="18q21.3"
<1..>911
/note="bcl2a mRNA"
147..764
/gene="BCL2"
147..764
/gene="BCL2"
/note="bcl2-beta protein"
/codon_start=1
/db_xref="GDB:G00-119-031"
/protein_id="AAA51814.1"
/db_xref="GI:179369"
/translation="MAHAGRTGYDNRREIVMKYIHYKLSQRYEWDAGDVGAAPPGAAP
APGIFSSQPGHTPHPAASRDPVARTPLQTPAAPGAAAGPALSVPVPVHLLALRQAGD
DFSRRYRGDFAEMSSQLHLTPFTARGFATVVEELFRDGVNWGRIVAFFEFGVMCVE

misc_feature 732
SVNREMSPLVDNIALWMTEYLNRLHHTWIQDNGGWVGASGDVSLG"
/gene="BCL2"
/note="alternative splice donor (intron A start)"
BASE COUNT 156 a 281 c 306 g 168 t
ORIGIN 556 bp upstream of SstI site.

Query Match 100.0%; Score 18; DB 67; Length 911;
Best Local Similarity 100.0%; Pred. No. 40;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 164 TCTCCAGCGTGCGCCAT 147

RESULT 15
AR054009/c
LOCUS AR054009 1846 bp DNA PAT 29-SEP-1999
DEFINITION Sequence 16 from patent US 5834306.
ACCESSION AR054009
VERSION AR054009.1 GI:5978871
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1846)
AUTHORS Webster,K.A. and Bishopric,N.H.
TITLE Tissue specific hypoxia regulated therapeutic constructs
JOURNAL Patent: US 5834306-A 16 10-NOV-1998;
FEATURES
source
1..1846
/organism="unknown"
BASE COUNT 424 a 520 c 483 g 419 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 1846;
Best Local Similarity 100.0%; Pred. No. 36;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgccat 18
|||||
Db 904 TCTCCAGCGTGCGCCAT 887

Search completed: December 4, 2000, 20:47:25
Job time: 18085 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 4, 2000, 16:32:02 ; Search time 99.31 Seconds
(without alignments)
68.089 Million cell updates/sec

Title: US-09-369-941-1
Perfect score: 18
Sequence: 1 tctcccagcgtgcgccat 18

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 480022 seqs, 187831343 residues

Total number of hits satisfying chosen parameters: 960044

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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21: /cgn2_2/gcgdata/geneseq/geneseq/NA2000.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	18	100.0	18	16 Q86659	Bcl-2 antisense oligo
2	18	100.0	18	19 V52545	Unmethylated CpG d
3	18	100.0	18	19 V27719	Immunostimulatory
4	18	100.0	18	19 V28181	Antisense oligonuc
5	18	100.0	18	19 V19667	Human bcl-2 antise
6	18	100.0	18	20 Z31944	CpG adjuvant oligo
7	18	100.0	18	20 Z41905	IL-12 secretion in
8	18	100.0	18	20 Z41948	IL-12 secretion in
9	18	100.0	18	20 X78803	HPV fusion protein
10	18	100.0	18	20 X88537	Cytosine-guanosine
11	18	100.0	18	20 X33514	BCL2-targeted anti
12	18	100.0	18	20 X23693	Deletion sequence

C	13	18	100.0	18	20	X27536	Synthetic RNA sequ
	14	18	100.0	18	20	X18702	Target bcl-2 antis
	15	18	100.0	18	20	V99434	Antisense oligonuc
	16	18	100.0	18	21	Z99003	CpG motif for immu
	17	18	100.0	18	21	Z60975	Nucleotide sequenc
	18	18	100.0	18	21	Z87997	BBTE-labeled oligo
	19	18	100.0	18	21	Z98660	Human Bcl-2 therap
	20	18	100.0	18	21	Z47643	Parasitic infectio
	21	18	100.0	18	21	Z47680	Parasitic infectio
	22	18	100.0	18	21	Z47850	Immunostimulatory
	23	18	100.0	18	21	Z47981	Immune remodeling
	24	18	100.0	18	21	Z48024	Immune remodeling
	25	18	100.0	20	20	V74246	CpG-N motif SOS-OD
	26	18	100.0	27	17	T18388	Human Bcl-2 forwar
C	27	18	100.0	35	16	Q86644	Bcl-2 translation
C	28	18	100.0	35	19	V19652	Human bcl-2 oligon
C	29	18	100.0	251	19	X11646	Human biallelic po
C	30	18	100.0	251	19	X12817	Human biallelic po
C	31	18	100.0	615	16	Q73987	Human bcl-2 gene O
C	32	18	100.0	760	17	T33694	Human BCL2 cDNA.
C	33	18	100.0	765	14	Q49815	Bcl-2. Homo sapie
C	34	18	100.0	831	9	N81293	Sequence of bcl-2
C	35	18	100.0	911	20	X08431	bcl-2 proto-oncoge
C	36	18	100.0	953	20	X33183	Bcl-2 DNA fragment
C	37	18	100.0	5086	15	Q54631	Human oncogene bcl
C	38	18	100.0	5086	16	Q86661	Human bcl-2 gene.
C	39	18	100.0	5086	19	X75766	Human bcl2 proto-o
C	40	18	100.0	5105	9	N81292	Sequence of bcl-2
C	41	18	100.0	7996	20	X33184	Base sequence of t
	42	17	94.4	17	19	V19659	Human bcl-2 antise
	43	16.4	91.1	18	19	V52559	Unmethylated CpG d
	44	16.4	91.1	18	19	V47681	Unmethylated CpG d
	45	16.4	91.1	18	19	V27733	Immunostimulatory

ALIGNMENTS

RESULT 1
Q86659
ID Q86659 standard; DNA; 18 BP.
XX
AC Q86659;
XX
DT 27-SEP-1995 (first entry)
XX
DE Bcl-2 antisense oligonucleotide.
XX
KW Anticodon oligomer; antisense oligonucleotide; bcl-2; cancer; therapy;
KW chemoresistance; ss.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT misc_feature 1..18
FT /*tag= a
FT /note= "3'-5' (antisense) sequence"
XX
PN WO9508350-A.
XX
PD 30-MAR-1995.
XX
PF 20-SEP-1994; 94WO-US10725.
XX
PR 20-SEP-1993; 93US-0124256.
XX
PA (REED/) REED J C.
XX
PI Reed JC;
XX
DR WPI; 1995-139394/18.
XX
PT Anti-code oligomers which bind to bcl-2 mRNA - for the treatment

PT of human solid tumours, esp. breast cancer
XX
PS Example 18; Page 44; 108pp; English.
XX
CC Reversal of chemoresistance of tumor cells by antisense-mediated
CC reduction of bcl1-2 expression was demonstrated using the
CC oligonucleotide given in Q86659. This is antisense to the first
CC 6 codons of the bcl-2 ORF.
XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 16; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctccagcgtgcgccat 18
Db 1 tctccagcgtgcgccat 18

RESULT 2
V52545
ID V52545 standard; DNA; 18 BP.
XX
AC V52545;
XX
DT 20-NOV-1998 (first entry)
XX
DE Unmethylated CpG dinucleotide 1758.
XX
KW Unmethylated CpG dinucleotide; immune response; bacterial meningitis;
KW natural killer cell activation; NK cell; Th2 response; neonatal sepsis;
KW pulmonary disorder; asthma; environmentally induced airway disease;
KW bacterial infection; endotoxaemia; therapy; cystic fibrosis;
KW inflammatory bowel disease; ss.
XX
OS Synthetic.
XX
PN WO9837919-A1.
XX
PD 03-SEP-1998.
XX
PF 25-FEB-1998; 98WO-US03678.
XX
PR 28-FEB-1997; 97US-0039405.
XX
PA (IOWA) UNIV IOWA RES FOUND.
XX
PI Krieg AM, Schwartz DA;
XX
DR WPI; 1998-480941/41.
XX
PT Use of nucleic acids containing an unmethylated CpG - for treating a
PT subject having or at risk of having an acute decrement in air flow
PT or inhibiting an inflammatory response
XX
PS Example 4; Page 35; 65pp; English.
XX
CC This sequence represents an unmethylated CpG dinucleotide, and can be
CC used in the method of the invention. The method is for treating a subject
CC having, or at risk of having an acute decrement in air flow, comprising
CC administering a nucleic acid sequence containing at least one
CC unmethylated CpG. The nucleic acids containing an unmethylated CpG
CC dinucleotide affect an immune response in a subject by activating natural
CC killer cells (NK) or redirecting a subject's immune response from a Th2
CC to a Th1 response by inducing monocytic and other cells to produce Th1
CC cytokines. They can be used to treat pulmonary disorders having an
CC immunologic component, such as asthma or environmentally induced airway
CC disease. They can also be used to treat diseases associated with
CC Gram-positive bacterial infections or endotoxaemia including bacterial
CC meningitis, neonatal sepsis, cystic fibrosis, inflammatory bowel disease
CC and liver cirrhosis, Gram-negative pneumonia, Gram-negative abdominal

CC abscess, haemorrhagic shock, disseminated intravascular coagulation, or
CC an inflammatory response to lipopolysaccharide.
XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 19; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctccagcgtgcgccat 18
Db 1 tctccagcgtgcgccat 18

RESULT 3
V27719
ID V27719 standard; DNA; 18 BP.
XX
AC V27719;
XX
DT 01-OCT-1998 (first entry)
XX
DE Immunostimulatory oligodeoxyribonucleotide of the invention.
XX
KW Immunostimulatory; oligodeoxyribonucleotide; ODN;
KW unmethylated CpG dinucleotide; activate; lymphocyte; immune response;
KW Th2; Th1; cytokine; treatment; prevention; asthma; autoimmune disease;
KW desensitisation therapy; artificial adjuvant; antibody generation; ss.
XX
OS Synthetic.
XX
PN WO9818810-A1.
XX
PD 07-MAY-1998.
XX
PF 30-OCT-1997; 97WO-US19791.
XX
PR 30-OCT-1996; 96US-0738652.
XX
PA (IOWA) UNIV IOWA RES FOUND.
XX
PI Kline JN, Krieg AM;
XX
DR WPI; 1998-272127/24.
XX
PT New immunostimulatory nucleic acid molecules - which contain at
PT least one unmethylated CpG dinucleotide, used for treating e.g.
PT tumours, infections or autoimmune disease
XX
PS Disclosure; Page 49; 109pp; English.
XX
CC V27641-751 represent immunostimulatory oligodeoxyribonucleotides (ODNs)
CC of the invention. The ODNs contain at least one unmethylated CpG
CC dinucleotide, and have the formula:
CC 5' N1X1CGX2N2 3', where at least one nucleotide separates consecutive
CC CpGs, X1 is adenine, guanine, or thymine, X2 is cytosine or thymine, N is
CC any nucleotide and N1+N2 is 0-26 bases with the provision that N1 and N2
CC does not contain a CCGG tetramer or more than one CCG or CGG trimer OR
CC 5' NX1X2CGX3X4N 3', where at least one nucleotide separates consecutive
CC CpGs, X1 and X2 are selected from CpT, GpG, GpA, ApT and ApA, X3and X4
CC are selected from TpT or CpT, N is any nucleotide and N1+N2 is 0-26
CC bases with the provision that N1 and N2 does not contain a CCGG tetramer
CC or more than one CCG or CGG trimer.
CC The ODNs activate lymphocytes in a subject and redirect a subject's
CC immune response from a Th2 to a Th1 (e.g. by inducing monocytic cells and
CC other cells to produce Th1 cytokines, including IL-12, IFN-gamma and
CC GM-CSF). The ODNs can be used to treat or prevent an asthmatic disorder,
CC autoimmune diseases, in desensitisation therapy, as an artificial
CC adjuvant during antibody generation in a mammal such as a mouse or a
CC human.
XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

```

Query Match      100.0%; Score 18; DB 19; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 tctcccagcgtgcgccat 18
Db      1 tctcccagcgtgcgccat 18

RESULT      4
V28181
ID      V28181 standard; DNA; 18 BP.
XX
AC      V28181;
XX
DT      08-OCT-1998 (first entry)
XX
DE      Antisense oligonucleotide to bcl-2 mRNA.
XX
KW      Purification; oligonucleotide; matrix; affinity unit;
KW      affinity purification; antisense; bcl-2; ss.
XX
OS      Synthetic.
XX
PN      WO9827425-A1.
XX
PD      25-JUN-1998.
XX
PF      18-DEC-1997; 97WO-US23284.
XX
PR      19-DEC-1996; 96US-0769951.
XX
PA      (ISIS-) ISIS PHARM INC.
XX
PI      Chen D, Cole DL, Srivatsa GS;
XX      WPI; 1998-362922/31.
XX
PT      Matrix for selective separation of oligo:nucleotide - useful for,
PT      e.g. large scale purification of anti-sense agents from their
PT      deletion derivatives formed during synthesis
XX
PS      Disclosure; Page 86; 183pp; English.
XX
CC      V28155-268 represent oligonucleotides which can be purified using the
CC      method of the invention. The specification describes a matrix that
CC      comprises a support and an affinity unit that specifically and reversibly
CC      binds a target oligonucleotide, and comprises a sequence of bases having
CC      the reverse complement of a hybridising portion of the target
CC      oligonucleotide. The matrix is used for affinity purification of
CC      synthetic oligonucleotides, specifically antisense agents, for treatment
CC      of hyperproliferative diseases, for treating a non-pathogen,
CC      non-hyperproliferative disease, e.g. Alzheimer's, for modulating
CC      expression of cell surface proteins, and to inhibit a eukaryotic
CC      pathogen, retrovirus or other viruses.
XX
SQ      Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match      100.0%; Score 18; DB 19; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 tctcccagcgtgcgccat 18
Db      1 tctcccagcgtgcgccat 18

RESULT      5
V19667
ID      V19667 standard; DNA; 18 BP.
```

```

XX      V19667;
AC
XX      12-JUN-1998 (first entry)
DT
XX      Human bcl-2 antisense oligonucleotide 13.
DE
XX      Antisense oligonucleotide; bcl-2 gene; lymphoma; leukaemia; human;
KW      cancer; ss.
KW
XX      Synthetic.
OS
OS      Homo sapiens.
XX
PN      US5734033-A.
XX
PD      31-MAR-1998.
XX
PF      24-MAR-1994; 94US-0288692.
XX
PR      21-FEB-1992; 92US-0840716.
PR      22-DEC-1988; 88US-0288692.
PR      24-MAR-1994; 94US-0217082.
XX
PA      (UYPE-) UNIV PENNSYLVANIA.
XX
PI      Reed J;
XX
XX      WPI; 1998-229881/20.
DR
XX
PT      Anti-sense oligo:nucleotide(s) complementary to BCL-2 mRNA - useful
PT      for treating cancers, e.g. lymphoma(s) and some leukaemia(s)
XX
PS      Disclosure; Column 23; 21pp; English.
XX
CC      This antisense oligonucleotide is complementary to the translation
CC      initiation site of the human bcl-2 mRNA. The Bcl-2 antisense
CC      oligonucleotides are phosphorothioate derivatives and can straddle
CC      strategic sites such as the translation initiation site, donor and
CC      acceptor splicing sites, or sites for transportation or degradation.
CC      Blocking translation at such strategic sites prevents the formation of
CC      a functional bcl-2 gene product. These oligonucleotides may be used for
CC      treating cancers associated with high levels of bcl-2 gene expression,
CC      especially lymphomas and some leukaemias.
XX
SQ      Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match      100.0%; Score 18; DB 19; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 tctcccagcgtgcgccat 18
Db      1 tctcccagcgtgcgccat 18

RESULT      6
Z31944
ID      Z31944 standard; DNA; 18 BP.
XX
AC      Z31944;
XX
DT      26-JAN-2000 (first entry)
XX
DE      CpG adjuvant oligo 1002.
XX
KW      CpG adjuvant; vaccine; polyoxyethylene ether; polyoxyethylene ester;
KW      antigen; infection; allergy; cancer; therapy; ss.
XX
OS      Synthetic.
XX
PN      WO9952549-A1.
XX
```


PD 21-OCT-1999.
XX
PF 29-MAR-1999; 99WO-EP02278.
XX
PR 09-APR-1998; 98GB-0007805.
PR 25-SEP-1998; 98GB-0020956.
XX
PA (SMIK) SMITHKLINE BEECHAM BIOLOGICALS.
XX
PI Friede M, Hermand P;
XX
DR WPI; 1999-620290/53.
XX
PT Vaccine to protect against infections, allergy and cancer -
XX
PS Example 9; Page 26; 52pp; English.
XX
CC This sequence represents a CpG adjuvant that can be used in the vaccine
CC composition of the invention. The vaccine comprises a polyoxyethylene
CC ether or ester (I), not in the form of a vesicle, pharmaceutically
CC acceptable excipient and an antigen (Ag) or antigenic composition. The
CC vaccine can be used to treat or prevent infections (by bacteria, viruses
CC or other parasites), allergy and cancer. (I), which are safe, easy to
CC sterilize and simple to administer, are powerful vaccine adjuvants, able
CC to induce a systemic immune response when administered (non-invasively)
CC to the mucosa. The response is at least as good as that from conventional
CC systemic injection. (I) are effective at low concentration, have low
CC reactogenicity and are well tolerated.
XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
Db ||||||
1 tctcccagcgtgcgccat 18

RESULT 7
241905
ID 241905 standard; DNA; 18 BP.
XX
AC 241905;
XX
DT 24-JAN-2000 (first entry)
XX
DE IL-12 secretion inducing CpG oligonucleotide 50.
XX
KW CpG oligonucleotide; phosphorothioate; interleukin-12; IL-12; secretion;
KW human PBMC; immune response; cancer; HIV; bacterial disease; asthma;
KW neoplastic disorder; jaagsiekte; B cell; NK cell; ss; cytokine;
KW antigen presenting cell; infection; allergic disease.
XX
OS Synthetic.
XX
PN WO9951259-A2.
XX
PD 14-OCT-1999.
XX
PF 02-APR-1999; 99WO-US07335.
XX
PR 03-APR-1998; 98US-0080729.
XX
PA (IOWA) UNIV IOWA RES FOUND.
XX
PI Krieg AM, Weiner G;
XX
DR WPI; 1999-620169/53.
XX
PT Novel synergistic combinations of immunostimulatory oligonucleotides

PT and immunopotentiating cytokines are useful for stimulating the immune
PT system -
XX
PS Example 8; Page 80; 91pp; English.
XX
CC Sequences 241856-241949 are phosphorothioate CpG oligonucleotides which
CC are used in the invention to induce interleukin-12 (IL-12) secretion
CC from human PBMC. The invention comprises stimulating an immune response
CC in a subject comprising administering to a subject exposed to an antigen,
CC an immunopotentiating cytokine and an immunostimulatory CpG
CC oligonucleotide to induce a synergistic antigen specific immune response.
CC The methods are useful for treating cancer by stimulating an antigen
CC specific immune response against a cancer antigen. The methods can also
CC be used to treat neoplastic disorders in humans, including but not
CC limited to: sarcoma, carcinoma, fibroma, lymphoma, melanoma,
CC neuroblastoma, retinoblastoma, and glioma. The methods are also useful
CC for treating infectious diseases, e.g. viral diseases such as HIV,
CC bacterial diseases, and fungal diseases. The methods may also be used to
CC treat allergic diseases, e.g. asthma. The methods and compositions may
CC also be applied to treat cancer and tumours in non human subjects,
CC e.g. cats and dogs. Neoplasias affecting agricultural livestock may also
CC be treated and include leukaemia, haemangiopericytoma and bovine ocular
CC neoplasia. Chronic, infectious, contagious diseases of sheep and goats
CC caused by the bacterium Corynebacterium pseudotuberculosis, and
CC contagious lung tumour of sheep caused by Jaagsiekte may also be treated.
CC CpG oligonucleotides can be useful in activating B cells, NK cells, and
CC antigen presenting cells, such as monocytes and macrophages. CpG
CC oligonucleotides enhance antibody dependent cellular cytotoxicity and can
CC be used as an adjuvant in conjunction with tumour antigens to protect
CC against a tumour challenge.
XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
Db ||||||
1 tctcccagcgtgcgccat 18

RESULT 8
241948
ID 241948 standard; DNA; 18 BP.
XX
AC 241948;
XX
DT 24-JAN-2000 (first entry)
XX
DE IL-12 secretion inducing CpG oligonucleotide 93.
XX
KW CpG oligonucleotide; phosphorothioate; interleukin-12; IL-12; secretion;
KW human PBMC; immune response; cancer; HIV; bacterial disease; asthma;
KW neoplastic disorder; jaagsiekte; B cell; NK cell; ss; cytokine;
KW antigen presenting cell; infection; allergic disease.
XX
OS Synthetic.
XX
PN WO9951259-A2.
XX
PD 14-OCT-1999.
XX
PF 02-APR-1999; 99WO-US07335.
XX
PR 03-APR-1998; 98US-0080729.
XX
PA (IOWA) UNIV IOWA RES FOUND.
XX
PI Krieg AM, Weiner G;
XX
DR WPI; 1999-620169/53.

XX Novel synergistic combinations of immunostimulatory oligonucleotides
PT and immunopotentiating cytokines are useful for stimulating the immune
PT system -
XX
XX Example 8; Page 88; 91pp; English.
PS
XX Sequences Z41856-Z41949 are phosphorothioate CpG oligonucleotides which
CC are used in the invention to induce interleukin-12 (IL-12) secretion
CC from human PBMC. The invention comprises stimulating an immune response
CC in a subject comprising administering to a subject exposed to an antigen,
CC an immunopotentiating cytokine and an immunostimulatory CpG
CC oligonucleotide to induce a synergistic antigen specific immune response.
CC The methods are useful for treating cancer by stimulating an antigen
CC specific immune response against a cancer antigen. The methods can also
CC be used to treat neoplastic disorders in humans, including but not
CC limited to: sarcoma, carcinoma, fibroma, lymphoma, melanoma,
CC neuroblastoma, retinoblastoma, and glioma. The methods are also useful
CC for treating infectious diseases, e.g. viral diseases such as HIV,
CC bacterial diseases, and fungal diseases. The methods may also be used to
CC treat allergic diseases, e.g. asthma. The methods and compositions may
CC also be applied to treat cancer and tumours in non human subjects,
CC e.g. cats and dogs. Neoplasias affecting agricultural livestock may also
CC be treated and include leukaemia, haemangiopericytoma and bovine ocular
CC neoplasia. Chronic, infectious, contagious diseases of sheep and goats
CC caused by the bacterium Corynebacterium pseudotuberculosis, and
CC CpG oligonucleotides can be useful in activating B cells, NK cells, and
CC antigen presenting cells, such as monocytes and macrophages. CpG
CC oligonucleotides enhance antibody dependent cellular cytotoxicity and can
CC be used as an adjuvant in conjunction with tumour antigens to protect
CC against a tumour challenge.
XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgccat 18
|||
Db 1 tctcccagcgtgcgccat 18

RESULT 9
X78803
ID X78803 standard; DNA; 18 BP.
XX
AC X78803;
XX
DT 06-SEP-1999 (first entry)
XX
DE HPV fusion protein CpG oligonucleotide 2.

XX Fusion protein; E6 protein; E7 protein; E6/E7; immunomodulator; tumour;
KW immunological fusion partner; CpG oligonucleotide; immune response;
KW HPV antigen; prevention; treatment; primer; ss.
XX

OS Synthetic.
OS Human papillomavirus.
XX
XX WO9933868-A2.
PN
XX
PD 08-JUL-1999.
XX
PF 18-DEC-1998; 98WO-EP08563.
XX
XX 24-DEC-1997; 97GB-0027262.
PR
XX (SMIK) SMITHKLINE BEECHAM BIOLOGICALS.
PA
XX Dalemans WLJ, Gerard CMG;

XX WPI; 1999-405485/34.
DR
XX Composition comprising an E6, E7 or E6/E7 fusion protein from HPV to
PT induce immune response to HPV
PT
XX
XX Claim 11; Page 37; 62pp; English.
PS
XX X78791-X78801 represent nucleic acid sequences which encode novel
CC constructs comprising an E6 or E7 protein or E6/E7 fusion protein from
CC HPV (represented in Y25375-Y25386). These constructs are optionally
CC linked to an immunological fusion partner and an immunomodulatory CpG
CC oligonucleotide. The products of the invention can be used to induce an
CC immune response in a patient to an HPV antigen. They can also be used
CC for preventing or treating HPV induced tumours. This sequence represents
CC a CpG oligonucleotide which is used in the method of the invention.
XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgccat 18
|||
Db 1 tctcccagcgtgcgccat 18

RESULT 10
X88537
ID X88537 standard; DNA; 18 BP.
XX
AC X88537;
XX
DT 10-SEP-1999 (first entry)
XX
DE Cytosine-guanosine dinucleotide motif oligonucleotide #4.

XX Cytosine-guanosine dinucleotide motif; CpG; immunomodulation;
KW unmethylated; vaccine; immunostimulation; immune response;
KW T-independent type 1 antigen; T-independent type 2 antigen;
KW polysaccharide conjugate antigen; ss.
XX
OS Synthetic.
XX
PN WO9933488-A2.
XX
PD 08-JUL-1999.
XX
PF 18-DEC-1998; 98WO-EP08562.
XX
PR 24-DEC-1997; 97GB-0027262.

XX (SMIK) SMITHKLINE BEECHAM BIOLOGICALS.
PA
XX Dalemans WLJ, Laferriere CAJ, Prieels J;
PI
XX WPI; 1999-405369/34.
DR
XX
XX A vaccine composition for inducing a immune response to
PT T-independent type 1 or type 2 antigen or polysaccharide conjugate
PT antigen
XX
PS Claim 6; Page 31; 35pp; English.
XX
CC The present invention describes a formulation (A) comprising a
CC cytosine-guanosine dinucleotide motif (CpG) oligonucleotide and
CC T-independent type 1 or type 2 antigens or polysaccharide conjugate
CC antigen. The present sequence represent a specifically claimed CpG
CC oligonucleotide. A vaccine composition comprising the formulation is
CC used for inducing a immune response to T-independent type 1 or type 2
CC antigen or polysaccharide conjugate antigen. The use of

CC immunostimulatory CpG oligonucleotide acts as an adjuvant to
CC pneumococcal polysaccharides.
XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 1 tctcccagcgtgcgccat 18

RESULT 11
X33514
ID X33514 standard; DNA; 18 BP.
XX
AC X33514;
XX
DT 07-JUL-1999 (first entry)
XX
DE BCL2-targeted antisense oligonucleotide SEQ ID NO:45.
XX
KW Combinatorial antisense library; oligonucleotide analogue; RNase;
KW ribozyme; cleavage; anchor; binding; target RNA; ss.
XX
OS Synthetic.
XX
PN WO9918238-A1.
XX
PD 15-APR-1999.
XX
PF 28-SEP-1998; 98WO-US20361.
XX
PR 18-AUG-1998; 98US-0136080.
PR 02-OCT-1997; 97US-0060673.
XX
PA (OASI-) OASIS BIOSCIENCES INC.
XX
PI Arnold LJ, Brown BD, Riley TA;
XX WPI; 1999-264039/22.
XX
PT Oligonucleotide analog compositions capable of coupling to form
PT antisense molecules
XX
PS Example 9; Page 45; 71pp; English.
XX

CC The present invention describes a composition comprising two
CC oligonucleotide analogues, each having a binding domain and a coupling
CC moiety, where the binding domains are capable of hybridizing to a target
CC polynucleotide and the coupling moieties are capable of coupling to each
CC other in the absence of a target molecule. The composition/compound is
CC used to cleave an RNA target. The compositions can be used to determine
CC an optimal antisense site for a given mRNA or an optimal ribozyme
CC cleavage site for a target RNA. By separating the antisense molecules
CC into two or more pieces, a comprehensive antisense library can be
CC prepared in advance, rather than synthesizing a plurality of candidate
CC antisense molecules as needed. A complete library of every possible
CC 17-mer oligonucleotide, using the four natural bases, would consist of
CC 417 (or about 1.7 x 1010) molecules. By providing the antisense molecules
CC in at least two components, e.g. a library of 8-mers and a library of
CC 9-mers, assembled quickly as needed, the library size is reduced to 48 +
CC 49, or 327 650 molecules. The complexity of the library can be further
CC reduced by substituting one or more universal or degenerate bases for
CC some of the natural bases. The present sequence represents an
CC oligonucleotide, which is used in an example from the present invention.
XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 1 tctcccagcgtgcgccat 18

RESULT 12
X23693
ID X23693 standard; DNA; 18 BP.
XX
AC X23693;
XX
DT 18-JUN-1999 (first entry)
XX
DE Deletion sequence oligonucleotide 146.
XX

KW Deletion sequence oligonucleotide; sensor array; eukaryotic pathogen;
KW probe; cellular adhesion modulator; cellular proliferation modulator;
KW human retrovirus; human immunodeficiency virus; non-human retrovirus;
KW HIV; primer; ss.
XX

OS Synthetic.
XX
PN WO9911820-A1.
XX
PD 11-MAR-1999.
XX

XX
PF 01-SEP-1998; 98WO-US18084.
XX

XX
PR 02-SEP-1997; 97US-0923771.
XX

XX
PA (ISIS-) ISIS PHARM INC.
XX

PI Chen D, Srivatsa GS;
XX

XX
DR WPI; 1999-205198/17.
XX

PT New compositions comprising sensor arrays made up of unique probe
PT oligonucleotides - useful for characterizing a sample of target
PT deletion oligonucleotides
XX

PS Example 9; Page 152; 163pp; English.
XX

CC This invention describes a novel composition comprising a number of
CC sensor arrays, where each array comprises a unique probe oligonucleotide,
CC which is the reverse complement of part of a unique target
CC oligonucleotide present in a mixture of target deletion sequence
CC oligonucleotides. The compositions form a method for characterizing a
CC sample of target deletion oligonucleotides which are labelled and
CC hybridize with the probe oligonucleotides of the sensor arrays. Such
CC oligonucleotides and their targets are represented in X23548-X23709.
CC Oligonucleotides characterized by the method form pharmaceutical
CC compositions that are useful for modulating cellular adhesion or
CC proliferation, and being active against a eukaryotic pathogen, a human
CC retrovirus, a human immunodeficiency virus (HIV), or a non-human
CC retrovirus, including influenza virus, Epstein-Barr virus, Respiratory
CC Syncytial Virus or cytomegalovirus (CMV). The compositions enable
CC characterization of deletion sequence oligonucleotides having related,
CC but different nucleobase sequences, and quantification of different
CC species of deletion sequence ("target") oligonucleotides in a mixture.
CC Also, if the specificity of the oligonucleotide's nucleobase sequence
CC for its reverse complement is not modified, the method may be performed
CC using oligodeoxynucleotides.
XX

SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
Db 1 tctcccagcgtgcgccat 18

RESULT 13
X27536/c
ID X27536 standard; RNA; 18 BP.
XX
AC X27536;
XX
DT 27-MAY-1999 (first entry)
XX
DE Synthetic RNA sequence produced by the method of the invention.
XX
KW Silyloxymethyl; phosphonate; silyloxymethyl halide; diagnosis; ss;
KW cyanoethyl phosphoramidate coupling; isomerisation; steric hindrance.
XX
OS Synthetic.
XX
PN WO9909044-A1.
XX
PD 25-FEB-1999.
XX
PF 17-AUG-1998; 98WO-EP05215.
XX
PR 18-AUG-1997; 97CH-0001931.
XX
PA (JENN/) JENNY L.
PA (PITS/) PITSCH S.
PA (WEIS/) WEISS P A.
XX
PI Jenny L, Pitsch S, Weiss PA;
PI WPI; 1999-180963/15.
DR
XX
PT 2-Silyloxymethyl ribonucleosides and their phosphonate derivatives
PT - have high purity, use in machine synthesis of ribonucleic acids,
PT enable longer oligonucleotide chain construction, and larger amounts
XX
PS Example 7; Page 26; 38pp; English.
XX
CC The invention relates to silyloxymethyl protected D- or L-ribonucleosides
CC and their phosphonates (I), and silyloxymethyl halides (II). (I) are
CC intermediates for synthesis of RNA-oligonucleotides with predetermined
CC nucleotide sequence, particularly by machine synthesis. The groups
CC specified above, apart from those on silyl, are those particularly for
CC the cyanoethyl phosphoramidate coupling. Uses of the oligoribonucleotide
CC products in diagnosis, therapy, and as research tools, are well known,
CC and are not dealt with in detail. (II) is an intermediate for (I). The
CC silyloxymethyl halide reagent is easy to prepare, and yields are high.
CC Introduction of the silyloxymethyl group into the ribonucleoside is
CC simple and rapid, and the acetal bond formed does not migrate,
CC eliminating particularly the prior art problem of 2' to 3' isomerisation.
CC The methylenedioxy group spacer between the silyl group and nucleoside
CC ring results in less steric hindrance than bulky direct silyloxy
CC linkages, enabling first, a range of choices for the silyl substituents,
CC to provide, e.g., acid or base stability; and second, higher yields in
CC coupling. Purer products are therefore obtained than in prior art,
CC enabling larger quantities and longer chains of oligoribonucleotides to
CC be synthesised successfully, and in shorter times.
XX
SQ Sequence 18 BP; 4 A; 4 C; 8 G; 2 U; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
Db 18 TCCTCCAGCGTGCGCCAT 1

RESULT 14
X18702
ID X18702 standard; DNA; 18 BP.
XX
AC X18702;
XX
DT 10-MAY-1999 (first entry)
XX
DE Target bcl-2 antisense oligonucleotide BCL-2.
XX
KW Cellular adhesion protein; proliferation; antisense oligonucleotide;
KW alimentary canal; transport; gastrointestinal mucosa; cancer;
KW Alzheimer's disease; beta-thalassemia; malaria; viral infection;
KW HIV; inflammation; ss.
XX
OS Synthetic.
XX
PN WO9901579-A1.
XX
PD 14-JAN-1999.
XX
PF 01-JUL-1998; 98WO-US13574.
XX
PR 01-JUL-1997; 97US-0886829.
XX
PA (ISIS-) ISIS PHARM INC.
XX
PI Hardee G, Teng C;
PI WPI; 1999-106077/09.
DR
XX
PT Composition comprising nucleic acid and penetration enhancer - used
PT particularly for delivering therapeutic antisense oligonucleotides
PT across the gastrointestinal mucosa, provides high bioavailability
XX
PS Example 2; Page 86; 115pp; English.
XX
CC A pharmaceutical composition has been developed which comprises a
CC nucleic acid and at least one penetration enhancer. The compositions are
CC used: (i) to treat or prevent any disease or disorder that can be
CC treated with the nucleic acid, e.g. cancer, Alzheimer's disease,
CC beta-thalassemia, malaria, viral infections (including human immune
CC deficiency virus (HIV)), inflammation, in human or animal medicine;
CC (ii) to investigate the role of a gene or gene product in non-human
CC animals; and (iii) to modulate gene expression in cells, tissues or
CC organs. The compositions provide bioavailability of at least 15,
CC preferably 17-35,%. The penetration enhancer improves: (i) transport of
CC the nucleic acid across the mucosa of the alimentary canal and into
CC cells; and (ii) increases stability of the nucleic acid. Oral
CC administration avoids the complications and expense of intravenous or
CC other methods of administration. X18669 to X18799 and X18801 represent
CC antisense oligonucleotides which can be used as the nucleic acid in
CC the method of the invention.
XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
Db 1 tctcccagcgtgcgccat 18

RESULT 15
V99434
ID V99434 standard; DNA; 18 BP.
XX
AC V99434;

```
XX 22-MAR-1999 (first entry)
XX Antisense oligonucleotide directed against human bcl-2 gene.
DE
XX Antisense oligonucleotide; human bcl-2 gene; phosphorothioate;
KW phosphodiester; lipid-encapsulation; tumour; aberrant gene expression;
KW treatment; inflammation; infection; ss.
XX
OS Synthetic.
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT modified_base 1..18
FT /*tag= a
FT /note= "phosphorothioate or phosphodiester bonds"
XX
PN WO9851278-A2.
XX
XX 19-NOV-1998.
XX
XX 14-MAY-1998; 98WO-CA00485.
XX
XX 14-MAY-1997; 97US-0856374.
XX
XX (INEX-) JINEX PHARM CORP.
XX
XX Ansell SM, Cullis P, Debeyer D, Harasym T, Hope MJ;
PI Klimuk SK, Scherrer P, Sempke SC;
XX
XX WPI; 1999-045179/04.
XX
XX Composition containing lipid-encapsulated therapeutic agent -
PT useful, e.g. for delivering antisense molecules or ribozymes or
PT treating diseases associated with aberrant gene expression
XX
PS Disclosure; Page 23; 98pp; English.
XX
CC The present sequence represents an antisense oligonucleotide directed
CC against the human bcl-2 gene. The oligonucleotide can have either
CC phosphorothioate or phosphodiester bonds. The oligonucleotide is
CC lipid-encapsulated using the method of the invention. A composition
CC comprising lipid-encapsulated particles of a therapeutic agent,
CC e.g. antisense oligonucleotides, is prepared by mixing at least
CC 2 lipids with buffered aqueous solution of charged therapeutic
CC agent to form an intermediate mixture of lipid-encapsulated particles,
CC and changing the pH of the mixture to neutralise at least some of the
CC external surface charges on the particles. One lipid has a
CC (de)protonatable group with Ka such that the lipid is charged at a
CC first pH but neutral at a second pH (particularly near physiological pH)
CC and the buffer maintains this lipid in the charged form (i.e. cationic
CC when the therapeutic agent is anionic in the buffer, or vice versa). The
CC second lipid prevents particle aggregation during formation of the
CC lipid-therapeutic agent particles. The composition is used to introduce
CC therapeutic agents into cells, in vivo or in vitro, particularly to
CC treat or prevent diseases associated with aberrant gene expression in
CC mammals, specifically tumours, inflammation or infection.
XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;
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Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 1 tctcccagcgtgcgccat 18
   |||
Db 1 tctcccagcgtgcgccat 18
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GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: December 4, 2000, 16:28:02 ; Search time 75.06 seconds
(without alignments)
36.269 Million cell updates/sec

Title: US-09-369-941-1
Perfect score: 18
Sequence: 1 tctcccagcgtgcgccat 18

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 262060 seqs, 75620727 residues

Total number of hits satisfying chosen parameters: 524120

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued_Patents_NA:*
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3: /cgn2_6/ptodata/2/ina/5C_COMB.seq.*
4: /cgn2_6/ptodata/2/ina/5D_COMB.seq.*
5: /cgn2_6/ptodata/2/ina/6_COMB.seq.*
6: /cgn2_6/ptodata/2/ina/PCTUS_COMB.seq.*
7: /cgn2_6/ptodata/2/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Query %	Length	DB	ID	Description
1	18	100.0	18	2	US-08-217-082A-17		Sequence 17, Appl
2	18	100.0	18	3	US-08-465-485A-17		Sequence 17, Appl
3	18	100.0	18	3	US-08-465-485A-24		Sequence 24, Appl
4	18	100.0	18	5	US-09-080-285-17		Sequence 17, Appl
5	18	100.0	18	5	US-09-080-285-24		Sequence 24, Appl
6	18	100.0	18	5	US-09-249-730-218		Sequence 218, App
7	18	100.0	27	1	US-08-410-804-13		Sequence 13, Appl
8	18	100.0	27	2	US-08-607-269-8		Sequence 8, Appli
9	18	100.0	27	2	US-08-259-514-13		Sequence 13, Appl
10	18	100.0	27	3	US-08-858-311-13		Sequence 13, Appl
11	18	100.0	27	6	PCT-US95-04600-8		Sequence 8, Appli
12	18	100.0	35	2	US-08-217-082A-2		Sequence 2, Appli
13	18	100.0	35	3	US-08-465-485A-2		Sequence 2, Appli
14	18	100.0	35	5	US-09-080-285-2		Sequence 2, Appli
15	18	100.0	615	3	US-08-465-485A-22		Sequence 22, Appl
16	18	100.0	615	5	US-09-080-285-22		Sequence 22, Appl
17	18	100.0	623	7	5506344-3		Patent No. 5506344
18	18	100.0	717	3	US-08-465-485A-20		Sequence 20, Appl
19	18	100.0	717	5	US-09-080-285-20		Sequence 20, Appl
20	18	100.0	760	2	US-08-405-702A-11		Sequence 11, Appl
21	18	100.0	831	7	5459251-3		Patent No. 5459251
22	18	100.0	831	7	5506344-4		Patent No. 5506344
23	18	100.0	911	6	PCT-US93-06251-3		Sequence 3, Appli
24	18	100.0	1846	3	US-08-365-486A-16		Sequence 16, Appl
25	18	100.0	4825	7	5459251-1		Patent No. 5459251
26	18	100.0	5086	3	US-08-465-485A-19		Sequence 19, Appl

C 27	18	100.0	5086	3	US-08-365-486A-14	Sequence 14, Appl
C 28	18	100.0	5086	5	US-09-080-285-19	Sequence 19, Appl
C 29	18	100.0	5086	6	PCT-US93-05651-4	Sequence 4, Appli
C 30	18	100.0	5086	6	PCT-US93-06251-2	Sequence 2, Appli
C 31	18	100.0	5104	7	5506344-1	Patent No. 5506344
C 32	17	94.4	17	2	US-08-217-082A-9	Sequence 9, Appli
C 33	15.4	85.6	33	5	US-08-650-726-1	Sequence 1, Appli
C 34	15	83.3	17	2	US-08-217-082A-8	Sequence 8, Appli
C 35	15	83.3	17	4	US-08-877-831-1	Sequence 1, Appli
C 36	14.8	82.2	335	7	5175102-1	Patent No. 5175102
C 37	14.4	80.0	6002	2	US-08-698-551-15	Sequence 15, Appl
C 38	14.4	80.0	6002	3	US-08-602-228-15	Sequence 15, Appl
C 39	14.4	80.0	6002	3	US-08-839-032A-15	Sequence 15, Appl
C 40	14.4	80.0	34303	4	US-08-735-609-4	Sequence 4, Appli
C 41	14.4	80.0	34303	4	US-08-735-609-4	Sequence 4, Appli
C 42	14.4	80.0	34303	5	US-09-315-372-4	Sequence 4, Appli
C 43	14.4	80.0	34303	5	US-09-244-752-4	Sequence 4, Appli
C 44	14.4	80.0	34303	5	US-09-245-497-4	Sequence 4, Appli
C 45	14.4	80.0	34382	3	US-08-374-483-6	Sequence 6, Appli

ALIGNMENTS

RESULT 1
US-08-217-082A-17
; Sequence 17, Application US/08217082A
; Patent No. 5734033
; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: ANTISENSE OLIGONUCLEOTIDES FOR INHIBITING THE
; TITLE OF INVENTION: GROWTH OF CELLS EXPRESSING THE HUMAN BCL-2 GENE
; NUMBER OF SEQUENCES: 17
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; ADDRESSEE: P.C.
; STREET: 224 Airport Parkway
; CITY: San Jose
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 95110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/217,082A
; FILING DATE: 24-MAR-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/840,716
; FILING DATE: 21-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/288,692
; FILING DATE: 22-DEC-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Fortney, Andrew D.
; REGISTRATION NUMBER: 34,600
; REFERENCE/DOCKET NUMBER: 3335-067-55 FWC
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (408) 436-2070
; TELEFAX: (408) 436-2075
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 18 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: Synthetic DNA

US-08-217-082A-17

Query Match 100.0%; Score 18; DB 2; Length 18;
Best Local Similarity 100.0%; Pred. No. 0.85;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
| | | | | | | | | | | | | | | | | |
Db 1 TCTCCAGCGTGCGCCAT 18

RESULT 2
US-08-465-485A-17
; Sequence 17, Application US/08465485A
; Patent No. 5831066
; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: Regulation of bcl-2 Gene Expression
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; ADDRESSEE: P.C.
; STREET: 1755 S. Jefferson Davis Hwy., Suite 400
; CITY: Arlington
; STATE: Virginia
; COUNTRY: U.S.A.
; ZIP: 22202

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/465,485A
; FILING DATE: 05-JUN-1995
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/124,256
; FILING DATE: 20-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/840,716
; FILING DATE: 21-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/288,692
; FILING DATE: 22-DEC-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Fortney, Andrew D.
; REGISTRATION NUMBER: 34,600
; REFERENCE/DOCKET NUMBER: 3335-070-55 CONT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (408) 436-2070
; TELEFAX: (408) 436-2075
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 18 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-465-485A-17

Query Match 100.0%; Score 18; DB 3; Length 18;
Best Local Similarity 100.0%; Pred. No. 0.85;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
| | | | | | | | | | | | | | | | | |
Db 1 TCTCCAGCGTGCGCCAT 18

RESULT 3
US-08-465-485A-24
; Sequence 24, Application US/08465485A
; Patent No. 5831066

; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: Regulation of bcl-2 Gene Expression
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; ADDRESSEE: P.C.
; STREET: 1755 S. Jefferson Davis Hwy., Suite 400
; CITY: Arlington
; STATE: Virginia
; COUNTRY: U.S.A.
; ZIP: 22202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/465,485A
; FILING DATE: 05-JUN-1995
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/124,256
; FILING DATE: 20-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/840,716
; FILING DATE: 21-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/288,692
; FILING DATE: 22-DEC-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Fortney, Andrew D.
; REGISTRATION NUMBER: 34,600
; REFERENCE/DOCKET NUMBER: 3335-070-55 CONT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (408) 436-2070
; TELEFAX: (408) 436-2075
; INFORMATION FOR SEQ ID NO: 24:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 18 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: Other nucleic acid;
; DESCRIPTION: Synthetic DNA
; ANTI-SENSE: YES
; FEATURE:
; NAME/KEY: Modified_base
; LOCATION: 16..17
; OTHER INFORMATION: Last two internucleoside linkages are
; OTHER INFORMATION: phosphorothioates
US-08-465-485A-24

Query Match 100.0%; Score 18; DB 3; Length 18;
Best Local Similarity 100.0%; Pred. No. 0.85;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
| | | | | | | | | | | | | | | | | |
Db 1 TCTCCAGCGTGCGCCAT 18

RESULT 4
US-09-080-285-17
; Sequence 17, Application US/09080285
; Patent No. 6040181
; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: Regulation of bcl-2 Gene Expression
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,

ADDRESSEE: P.C.
STREET: 1755 S. Jefferson Davis Hwy., Suite 400
CITY: Arlington
STATE: Virginia
COUNTRY: U.S.A.
ZIP: 22202
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/080,285
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/465,485
FILING DATE: 05-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/124,256
FILING DATE: 20-SEP-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/840,716
FILING DATE: 21-FEB-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/288,692
FILING DATE: 22-DEC-1988
ATTORNEY/AGENT INFORMATION:
NAME: Fortney, Andrew D.
REGISTRATION NUMBER: 34,600
REFERENCE/DOCKET NUMBER: 3335-070-55 CONT
TELEPHONE: (408) 436-2070
TELEFAX: (408) 436-2075
INFORMATION FOR SEQ ID NO: 17:
SEQUENCE CHARACTERISTICS:
LENGTH: 18 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-09-080-285-17

Query Match 100.0%; Score 18; DB 5; Length 18;
Best Local Similarity 100.0%; Pred. No. 0.85;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgccat 18
| | | | | | | | | | | | | | | |
Db 1 TCTCCCAGCGTGCGCCAT 18

RESULT 5
US-09-080-285-24
; Sequence 24, Application US/09080285
; Patent No. 6040181
; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: Regulation of bcl-2 Gene Expression
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; ADDRESSEE: P.C.
; STREET: 1755 S. Jefferson Davis Hwy., Suite 400
; CITY: Arlington
; STATE: Virginia
; COUNTRY: U.S.A.
; ZIP: 22202
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/080,285
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/465,485
FILING DATE: 05-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/124,256
FILING DATE: 20-SEP-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/840,716
FILING DATE: 21-FEB-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/288,692
FILING DATE: 22-DEC-1988
ATTORNEY/AGENT INFORMATION:
NAME: Fortney, Andrew D.
REGISTRATION NUMBER: 34,600
REFERENCE/DOCKET NUMBER: 3335-070-55 CONT
TELEPHONE: (408) 436-2070
TELEFAX: (408) 436-2075
INFORMATION FOR SEQ ID NO: 24:
SEQUENCE CHARACTERISTICS:
LENGTH: 18 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: Other nucleic acid;
DESCRIPTION: Synthetic DNA
ANTI-SENSE: YES
FEATURE:
NAME/KEY: Modified_base
LOCATION: 16..17
OTHER INFORMATION: Last two internucleoside linkages are
OTHER INFORMATION: phosphorothioates
US-09-080-285-24

Query Match 100.0%; Score 18; DB 5; Length 18;
Best Local Similarity 100.0%; Pred. No. 0.85;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgccat 18
| | | | | | | | | | | | | | | |
Db 1 TCTCCCAGCGTGCGCCAT 18

RESULT 6
US-09-249-730-218
; Sequence 218, Application US/09249730
; Patent No. 6121000
; GENERAL INFORMATION:
; APPLICANT: WRIGHT, Jim A.
; APPLICANT: YOUNG, Aiping H.
; TITLE OF INVENTION: Antitumor Antisense Sequences Directed Against R1 and
; TITLE OF INVENTION: R2 Components of Ribonucleotide Reductase
; FILE REFERENCE: 032396-040
; CURRENT APPLICATION NUMBER: US/09/249,730
; CURRENT FILING DATE: 1999-02-11
; NUMBER OF SEQ ID NOS: 220
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 218
; LENGTH: 18
; TYPE: DNA
; ORGANISM: Human
US-09-249-730-218

Query Match 100.0%; Score 18; DB 5; Length 18;
Best Local Similarity 100.0%; Pred. No. 0.85;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccaat 18
|||||
Db 1 tctcccagcgtgcgccaat 18

RESULT 7
US-08-410-804-13/c
; Sequence 13, Application US/08410804
; Patent No. 5632994
; GENERAL INFORMATION:
; APPLICANT: Reed, John C.
; APPLICANT: Sato, Takaaki
; TITLE OF INVENTION: FAS ASSOCIATED PROTEINS
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Cathryn Campbell
; STREET: 4370 La Jolla Village Drive. Ste 700
; CITY: San Diego
; STATE: California
; COUNTRY: United States
; ZIP: 92122
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/410,804
; FILING DATE: 27-MAR-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/259,514
; FILING DATE: 14-JUN-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Campbell, Cathryn
; REGISTRATION NUMBER: 31,815
; REFERENCE/DOCKET NUMBER: P-LJ 1389
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 535-9001
; TELEFAX: (619) 535-8949
; INFORMATION FOR SEQ ID NO: 13:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 27 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
US-08-410-804-13

Query Match 100.0%; Score 18; DB 1; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.86;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccaat 18
|||||
Db 25 TCTCCCAGCGTGCGCCAT 8

RESULT 8
US-08-607-269-8/c
; Sequence 8, Application US/08607269
; Patent No. 5702897
; GENERAL INFORMATION:
; APPLICANT: Reed, John C.
; APPLICANT: Sato, Takaaki
; TITLE OF INVENTION: Interaction of Proteins Involved in a
; TITLE OF INVENTION: Cell Death Pathway
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Campbell and Flores
; STREET: 4370 La Jolla Village Drive, Suite 700

; CITY: San Diego
; STATE: California
; COUNTRY: USA
; ZIP: 92122
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/607,269
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/226,876
; FILING DATE: 13-APR-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Campbell, Cathryn A.
; REGISTRATION NUMBER: 31,815
; REFERENCE/DOCKET NUMBER: P-LJ 9882
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 535-9001
; TELEFAX: (619) 535-8949
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 27 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-607-269-8

Query Match 100.0%; Score 18; DB 2; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.86;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccaat 18
|||||
Db 25 TCTCCCAGCGTGCGCCAT 8

RESULT 9
US-08-259-514-13/c
; Sequence 13, Application US/08259514
; Patent No. 5747245
; GENERAL INFORMATION:
; APPLICANT: Reed, John C.
; APPLICANT: Sato, Takaaki
; TITLE OF INVENTION: FAS ASSOCIATED PROTEINS
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Cathryn Campbell
; STREET: 4370 La Jolla Village Drive. Ste 700
; CITY: San Diego
; STATE: California
; COUNTRY: United States
; ZIP: 92122
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/259,514
; FILING DATE: 14-JUN-1994
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Campbell, Cathryn
; REGISTRATION NUMBER: 31,815
; REFERENCE/DOCKET NUMBER: P-LJ 9954
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 535-9001
; TELEFAX: (619) 535-8949

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; INFORMATION FOR SEQ ID NO: 13:
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; SEQUENCE CHARACTERISTICS:
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; LENGTH: 27 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: CDNA
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; US-08-259-514-13

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Query Match      100.0%; Score 18; DB 2; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.86;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Qy 1 tctccagcgtgcgccaat 18
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 Db 25 TCTCCCAGCGTGCGCCAT 8

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RESULT 10
US-08-858-311-13/c
; Sequence 13, Application US/08858311
; Patent No. 5876939
; GENERAL INFORMATION:
; APPLICANT: Reed, John C.
; APPLICANT: Sato, Takaaki
; TITLE OF INVENTION: FAS ASSOCIATED PROTEINS
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Cathryn Campbell
; STREET: 4370 La Jolla Village Drive. Ste 700
; CITY: San Diego
; STATE: California
; COUNTRY: United States
; ZIP: 92122
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/858,311
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Query Match      100.0%; Score 18; DB 3; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.86;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Qy 1 tctccagcgtgcgccat 18
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 pb 25 TCTCCCAGCGTGCGCCAT 8

RESULT 11
PCT-US95-04600-8/c
; Sequence 8, Application PC/TUS9504600
; GENERAL INFORMATION:
; APPLICANT: LA JOLLA CANCER RESEARCH FOUNDATION
; TITLE OF INVENTION: Interaction of Proteins Involved in
; TITLE OF INVENTION: a Cell Death Pathway
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Campbell and Flores
; STREET: 4370 La Jolla Village Drive, Suite 700
; CITY: San Diego
; STATE: California
; COUNTRY: USA
; ZIP: 92122
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/04600
; FILING DATE: 12-APR-1995
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Imbra, Richard J.
; REGISTRATION NUMBER: 37,643
; REFERENCE/DOCKET NUMBER: FP-LJ 1361
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 535-9001
; TELEFAX: (619) 535-8949
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 27 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; PCT-US95-04600-8

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Query Match      100.0%; Score 18; DB 6; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.86;
Matches 18: Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Qy 1 tctccagcgtgcgccat 18
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Db 25 TCTCCCAGCGTGGGCCAT 8

RESULT 12
US-08-217-082A-2/c
; Sequence 2, Application US/08217082A
; Patent No. 5734033
; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: ANTISENSE OLIGONUCLEOTIDES FOR INHIBITING THE
; TITLE OF INVENTION: GROWTH OF CELLS EXPRESSING THE HUMAN BCL-2 GENE
; NUMBER OF SEQUENCES: 17
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; ADDRESSEE: P.C.
; STREET: 224 Airport Parkway
; CITY: San Jose
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 95110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25

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; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/217,082A
; FILING DATE: 24-MAR-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/840,716
; FILING DATE: 21-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/288,692
; FILING DATE: 22-DEC-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Fortney, Andrew D.
; REGISTRATION NUMBER: 34,600
; REFERENCE/DOCKET NUMBER: 3335-067-55 FWC
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (408) 436-2070
; TELEFAX: (408) 436-2075
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 35 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; ANTI-SENSE: NO
US-08-217-082A-2

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Best Local Similarity 100.0%; Pred. No. 0.86;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 35 TCTCCCAGCGTGCGCCAT 18

RESULT 13

US-08-465-485A-2/c
; Sequence 2, Application US/08465485A
; Patent No. 5831066
; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: Regulation of bcl-2 Gene Expression
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; ADDRESSEE: P.C.
; STREET: 1755 S. Jefferson Davis Hwy., Suite 400
; CITY: Arlington
; STATE: Virginia
; COUNTRY: U.S.A.
; ZIP: 22202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/465,485A
; FILING DATE: 05-JUN-1995
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/124,256
; FILING DATE: 20-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/840,716
; FILING DATE: 21-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/288,692
; FILING DATE: 22-DEC-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Fortney, Andrew D.

;
;
;
; REGISTRATION NUMBER: 34,600
; REFERENCE/DOCKET NUMBER: 3335-070-55 CONT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (408) 436-2070
; TELEFAX: (408) 436-2075
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 35 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; ANTI-SENSE: NO
US-08-465-485A-2

Query Match 100.0%; Score 18; DB 3; Length 35;
Best Local Similarity 100.0%; Pred. No. 0.86;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 35 TCTCCCAGCGTGCGCCAT 18

RESULT 14

US-09-080-285-2/c
; Sequence 2, Application US/09080285
; Patent No. 6040181
; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: Regulation of bcl-2 Gene Expression
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; ADDRESSEE: P.C.
; STREET: 1755 S. Jefferson Davis Hwy., Suite 400
; CITY: Arlington
; STATE: Virginia
; COUNTRY: U.S.A.
; ZIP: 22202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/080,285
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/465,485
; FILING DATE: 05-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/124,256
; FILING DATE: 20-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/840,716
; FILING DATE: 21-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/288,692
; FILING DATE: 22-DEC-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Fortney, Andrew D.
; REGISTRATION NUMBER: 34,600
; REFERENCE/DOCKET NUMBER: 3335-070-55 CONT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (408) 436-2070
; TELEFAX: (408) 436-2075
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 35 base pairs
; TYPE: nucleic acid

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; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; - ANTI-SENSE: NO
US-09-080-285-2

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Query Match 100.0%; Score 18; DB 5; Length 35;
Best Local Similarity 100.0%; Pred. No. 0.86;
Matches 18; Conservative 0; Mismatches 0; Indels

Qy 1 tctcccagcgtgcgccat 18
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 Db 35 TCTCCCAGCGTGCGCCAT 18

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RESULT 15
US-08-465-485A-22/c
; Sequence 22, Application US/08465485A
; Patent No. 5831066
; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: Regulation of bcl-2 Gene Expression
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; ADDRESSEE: P.C.
; STREET: 1755 S. Jefferson Davis Hwy., Suite 400
; CITY: Arlington
; STATE: Virginia
; COUNTRY: U.S.A.
; ZIP: 22202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
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; FILING DATE: 05-JUN-1995
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/124,256
; FILING DATE: 20-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/840,716
; FILING DATE: 21-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/288,692
; FILING DATE: 22-DEC-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Fortney, Andrew D.
; REGISTRATION NUMBER: 34,600
; REFERENCE/DOCKET NUMBER: 3335-070-55 CONT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (408) 436-2070
; TELEFAX: (408) 436-2075
; INFORMATION FOR SEQ ID NO: 22:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 615 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..615
; US-08-465-485A-22

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Qy 1 tctcccagcgtgcgccat 18
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Db 18 TCTCCCAGCGTGCGCCAT 1

Search completed: December 4, 2000, 21:08:04
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Query Match          100.0%; Score 18; DB 3; Length 615;
Best Local Similarity 100.0%; Pred. No. 0.9;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Query Match 100.0%; Score 18; DB 3; Length 615;

Best Local Similarity 100.0%; Pred. No. 0.9;

Matches	18;	Conservative	0;	Mismatches	0;	Indels	0;	Gaps	0;
---------	-----	--------------	----	------------	----	--------	----	------	----

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 4, 2000, 21:06:39 ; Search time 1141.84 Seconds
(without alignments)
108.295 Million cell updates/sec

Title: US-09-369-941-2
Perfect score: 20
Sequence: 1 tccatgacgttcctgacgtt 20

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 7189864 seqs, 3091403243 residues

Total number of hits satisfying chosen parameters: 14379728

Minimum DB seq length: 0
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Post-processing: Minimum Match 0%
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Listing first 45 summaries

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126: em_gss15:*
127: em_gss16:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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4	16.8	84.0	821	114	CNS03G84	AL242653 Tetraodon
5	16.8	84.0	992	114	CNS040Q4	AL269221 Tetraodon
6	16.8	84.0	994	114	CNS0421L	AL271542 Tetraodon
7	16.4	82.0	824	113	CNS01XP2	AL172016 Tetraodon
8	16.4	82.0	1015	115	CNS05NA3	AL345108 Tetraodon
9	16	80.0	179	19	AV525865	AV525865 AV525865
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11	15.8	79.0	262	30	BB398655	BB398655 BB398655
12	15.8	79.0	392	8	AI077366	AI077366 oy87e11.x
13	15.8	79.0	408	8	AI086210	AI086210 ow90d05.s
14	15.8	79.0	490	19	AW063142	AW063142 TN0287 KR
15	15.8	79.0	509	98	AQ478992	AQ478992 RPCI-11-2
16	15.8	79.0	717	114	CNS04SXC	AL305773 Tetraodon
17	15.8	79.0	934	99	AQ573722	AQ573722 nbxb0084B
18	15.8	79.0	938	114	CNS03144	AL245101 Tetraodon
19	15.8	79.0	992	114	CNS04Q70	AL302229 Tetraodon
20	15.4	77.0	219	115	FR0021650	AL014523 F.rubripe
21	15.4	77.0	279	100	AQ651891	AQ651891 Sheared D
22	15.4	77.0	302	115	FR0023000	AL015860 F.rubripe
23	15.4	77.0	311	28	BB234999	BB234999 BB234999
24	15.4	77.0	317	115	FR0021653	AL014526 F.rubripe
25	15.4	77.0	347	115	FR0023012	AL015871 F.rubripe
26	15.4	77.0	395	115	FR0021632	AL014505 F.rubripe
27	15.4	77.0	467	9	AI181400	AI181400 uc59c05.r
28	15.4	77.0	537	13	AI812316	AI812316 10G3 Pine
29	15.4	77.0	552	115	FR0021654	AL014527 F.rubripe
30	15.4	77.0	554	115	FR0021629	AL014502 F.rubripe
31	15.4	77.0	580	115	FR0023009	AL015869 F.rubripe
32	15.4	77.0	584	115	FR0021635	AL014508 F.rubripe
33	15.4	77.0	597	115	FR0021641	AL014514 F.rubripe
34	15.4	77.0	598	115	FR0021637	AL014510 F.rubripe
35	15.4	77.0	609	115	FR0022992	AL015852 F.rubripe
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37	15.4	77.0	629	34	BE195266	BE195266 HVSMEH008
38	15.4	77.0	658	9	AI234463	AI234463 EST366 Ma
39	15.2	76.0	151	4	AA494709	AA494709 fa10h06.r
40	15.2	76.0	189	12	AI771817	AI771817 EST252917
41	15.2	76.0	253	30	BB385734	BB385734 BB385734
42	15.2	76.0	272	7	AA958394	AA958394 ua11c12.r
43	15.2	76.0	273	89	AQ105312	AQ105312 HS_3020_B
44	15.2	76.0	277	27	BB110967	BB110967 BB110967
45	15.2	76.0	278	28	BB230155	BB230155 BB230155

ALIGNMENTS

RESULT 1
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DEFINITION	zp24f01.s1 Stratagene neuroepithelium (#937231) Homo sapiens cDNA clone IMAGE:610393 3' similar to contains Alu repetitive element;; mRNA sequence.				
ACCESSION	AA171941				
VERSION	AA171941.1	GI:1751000			
KEYWORDS	EST.				
SOURCE	human.				
ORGANISM	Homo sapiens				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.				
AUTHORS	1 (bases 1 to 464) Hillier,L., Lennon,G., Becker,M., Bonaldo,M.F., Chiapelli,B., Chissoe,S., Dietrich,N., Dubuque,T., Favello,A., Gish,W., Hawkins,M., Hultman,M., Kucaba,T., Lacy,M., Le,M., Le,N., Mardis,E., Moore,B., Morris,M., Parsons,J., Prange,C., Rifkin,L., Rohlfing,T., Schellenberg,K., Soares,M.B., Tan,F., Thierry-Mieg,J., Trevaskis,E., Underwood,K., Wohlmann,P., Waterston,R., Wilson,R. and Marra,M. Generation and analysis of 280,000 human expressed sequence tags Genome Res. 6 (9), 807-828 (1996)				
TITLE	Genome Res. 6 (9), 807-828 (1996)				
JOURNAL	97044478				
MEDLINE	Contact: Wilson RK				
COMMENT	Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108 Tel: 314 286 1800 Fax: 314 286 1810 Email: estewatson.wustl.edu This clone is available royalty-free through LNL; contact the IMAGE Consortium (info@image.lnl.gov) for further information. Seq primer: -40M13 fwd. from Amersham High quality sequence stop: 360.				
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	/db_xref="taxon:9606"				
	/clone="IMAGE:610393"				
	/clone_lib="Stratagene neuroepithelium (#937231)"				
	/dev_stage="Ntera-2/RA neuroepithelial cells"				
	/lab_host="SOLR (kanamycin resistant)"				
	/note="Vector: pBluescript SK-; Site_1: EcoRI; Site_2: XhoI; Cloned unidirectionally. Primer: Oligo dt. NT2 cells (Ntera-2/cl.D1) induced with Retinoic Acid for 24 hours. Average insert size: 1.5 kb; Uni-ZAP XR Vector; ~5' adaptor sequence: 5' GAATTCGCACGAG 3' ~3' adaptor sequence: 5' CTCGAGTTTTTTTTTTTTTTT 3'"				
BASE COUNT	109 a	105 c	100 g	148 t	2 others
ORIGIN					
Query Match	84.0%; Score 16.8; DB 2; Length 464;				
Best Local Similarity	90.0%; Pred. No. 2.3e+02;				
Matches	18;	Conservative	0;	Mismatches	2; Indels 0; Gaps 0;
QY	1	ttcatgacgttcctgacgtt	20		
Db	431	TCCATGAGGTTCTCGAAGTT	450		
RESULT	2				
AW065908/c	546 bp	mRNA	EST	30-MAR-2000	
LOCUS	687002G08.y1	687	- Early embryo from Delaware Zea mays cDNA, mRNA sequence.		
DEFINITION	sequence.				
ACCESSION	AW065908				
VERSION	AW065908.1	GI:6020980			
KEYWORDS	EST.				
SOURCE	Zea mays.				
ORGANISM	Zea mays				
REFERENCE	Eukaryota; Viridiplantae; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Zea.				
AUTHORS	1 (bases 1 to 546) Walbot,V.				

TITLE Maize ESTs from various cDNA libraries sequenced at Stanford University
JOURNAL Unpublished (1999)
COMMENT Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Plate: 687002 row: G column: 08.
Location/Qualifiers
1. .546
/organism="Zea mays"
/cultivar="Illinois High Oil"
/db_xref="taxon:4577"
/clone_lib="687 - Early embryo from Delaware"
/tissue_type="embryo"
/dev_stage="14, 21, 28, and 35 days after pollination"
/lab_host="E. coli SOLR"
/note="Organ: embryo; Vector: pBluescript SK; Site_1: XhoI
; Site_2: EcoRI; Library was prepared by Statagene using
the Uni-ZAP XR system (Stratagene BN937328-12). Clones
were picked by a Q-bot after blue/white selection
(ampicillin resistance - use 100 micrograms/microliter).
Developed from a pool of equal amounts of RNA from
developing embryos sampled at 14, 21, 28 and 35 days after
pollination of the Illinois High Oil Maize Strain Cycle
90. This closed strain has been selected for high oil
concentration for 90 generations and originates from the
1890s era open pollinated variety Burr's White"

BASE COUNT 113 a 183 c 156 g 94 t
ORIGIN

Query Match 84.0%; Score 16.8; DB 19; Length 546;
Best Local Similarity 90.0%; Pred. No. 2.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 tccatgacgttcctgacgtt 20
|||||
Db 229 TCCATGACGTTCCTGATCTT 210

RESULT 3
CNS02N06 797 bp DNA GSS 14-MAY-2000
LOCUS Tetraodon nigroviridis genome survey sequence T7 end of clone
DEFINITION 151L08 of library G from Tetraodon nigroviridis, genomic survey
sequence.
ACCESSION AL205647
VERSION AL205647.1 GI:7864466
KEYWORDS GSS; genome survey sequence.
SOURCE Tetraodon nigroviridis.
ORGANISM Tetraodon nigroviridis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Eurypterygii; Ctenosquamata; Acanthomorpha; Euacanthomorpha;
Holacanthopterygii; Acanthopterygii; Percomorpha;
Tetraodontiformes; Tetraodontidae; Tetraodon.
REFERENCE 1 (bases 1 to 797)
AUTHORS Roest-Crollius,H., Jalllon,O., Dasilva,C., Fizames,C., Fisher,C.,
Bonneau,L., Billault,A., Quetier,F., Saurin,W., Bernot,A. and
Weissenbach,J.
TITLE Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 797)
AUTHORS Roest-Crollius,H., Jalllon,O., Dasilva,C., Bouneau,L., Fisher,C.,
Bernot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,
Saurin,W. and Weissenbach,J.
TITLE Human gene number estimate provided by genome wide analysis using
Tetraodon nigroviridis DNA sequence

JOURNAL Unpublished
REFERENCE 3 (bases 1 to 797)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (12-APR-2000) to the EMBL/GenBank/DBJ databases
COMMENT This sequence is a single read and was generated as part of a large
scale clone-end sequencing project of the Tetraodon nigroviridis
genome. For more information, please take a look at
<http://www.genoscope.cns.fr/Tetraodon>.
Location/Qualifiers
1. .797
/organism="Tetraodon nigroviridis"
/db_xref="taxon:99883"
/clone="151L08"
/clone_lib="G"
/note="Genoscope sequence ID : C0AG151DF04LPI-end : T7"

BASE COUNT 192 a 204 c 209 g 177 t 15 others
ORIGIN

Query Match 84.0%; Score 16.8; DB 113; Length 797;
Best Local Similarity 90.0%; Pred. No. 2.5e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 tccatgacgttcctgacgtt 20
|||||
Db 604 TCCAAAGACGTTCCTGGCGTT 585

RESULT 4
CNS03G84 821 bp DNA GSS 17-MAY-2000
LOCUS Tetraodon nigroviridis genome survey sequence T7 end of clone
DEFINITION 023N02 of library G from Tetraodon nigroviridis, genomic survey
sequence.
ACCESSION AL242653
VERSION AL242653.1 GI:7963422
KEYWORDS GSS; genome survey sequence.
SOURCE Tetraodon nigroviridis.
ORGANISM Tetraodon nigroviridis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Eurypterygii; Ctenosquamata; Acanthomorpha; Euacanthomorpha;
Holacanthopterygii; Acanthopterygii; Percomorpha;
Tetraodontiformes; Tetraodontidae; Tetraodon.
REFERENCE 1 (bases 1 to 821)
AUTHORS Roest-Crollius,H., Jalllon,O., Dasilva,C., Fizames,C., Fisher,C.,
Bonneau,L., Billault,A., Quetier,F., Saurin,W., Bernot,A. and
Weissenbach,J.
TITLE Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 821)
AUTHORS Roest-Crollius,H., Jalllon,O., Dasilva,C., Bouneau,L., Fisher,C.,
Bernot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,
Saurin,W. and Weissenbach,J.
TITLE Human gene number estimate provided by genome wide analysis using
Tetraodon nigroviridis DNA sequence
JOURNAL Unpublished
REFERENCE 3 (bases 1 to 821)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (12-APR-2000) to the EMBL/GenBank/DBJ databases
COMMENT This sequence is a single read and was generated as part of a large
scale clone-end sequencing project of the Tetraodon nigroviridis
genome. For more information, please take a look at
<http://www.genoscope.cns.fr/Tetraodon>.
Location/Qualifiers
1. .821
/organism="Tetraodon nigroviridis"
/db_xref="taxon:99883"
/clone="023N02"
/clone_lib="G"

BASE COUNT 217 a 145 c 172 g 279 t 8 others
ORIGIN

Query Match 84.0%; Score 16.8; DB 114; length 821;
Best Local Similarity 90.0%; Pred. No. 2.5e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
 |||||
Db 627 TCCATTGACGTTACTGACGTT 646

RESULT 5
CNS04004 992 bp DNA GSS 18-MAY-2000
LOCUS
DEFINITION Tetraodon nigroviridis genome survey sequence T7 end of clone
 073A13 of library G from Tetraodon nigroviridis, genomic survey
 sequence.

ACCESSION AL269221 GI:7991098
VERSION
KEYWORDS GSS: genome survey sequence.
SOURCE Tetraodon nigroviridis.
ORGANISM Tetraodon nigroviridis
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
 Eurypterygii; Ctenosquamata; Acanthomorpha; Euacanthomorpha;
 Holacanthopterygii; Acanthopterygii; Percomorpha;
 Tetraodontiformes; Tetraodontidae; Tetraodon.

REFERENCE 1 (bases 1 to 992)
AUTHORS Roest-Crollius,H., Jalllon,O., Dasilva,C., Fizames,C., Fisher,C.,
 Bouneau,L., Billault,A., Quetlier,F., Saurin,W., Bernot,A. and
 Weissenbach,J.
TITLE Characterization and repeat analysis of the compact genome of the
 freshwater pufferfish Tetraodon nigroviridis

JOURNAL Unpublished
REFERENCE 2 (bases 1 to 992)
AUTHORS Roest-Crollius,H., Jalllon,O., Dasilva,C., Bouneau,L., Fisher,C.,
 Bernot,A., Fizames,C., Wincker,P., Brotlier,P., Quetlier,F.,
 Saurin,W. and Weissenbach,J.
TITLE Human gene number estimate provided by genome wide analysis using
 Tetraodon nigroviridis DNA sequence
JOURNAL Unpublished
REFERENCE 3 (bases 1 to 992)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (12-APR-2000) to the EMBL/GenBank/DBJ databases
COMMENT This sequence is a single read and was generated as part of a large
 scale clone-end sequencing project of the Tetraodon nigroviridis
 genome. For more information, please take a look at
 http://www.genoscope.cns.fr/Tetraodon.

FEATURES
source
1..992
/organism="Tetraodon nigroviridis"
/db_xref="taxon:99883"
/clone="073A13"
/clone_1lb="G"
/note="Genoscope sequence ID : C0BG073AA07LP1-end : T7"

BASE COUNT 242 a 184 c 211 g 337 t 18 others
ORIGIN

Query Match 84.0%; Score 16.8; DB 114; length 992;
Best Local Similarity 90.0%; Pred. No. 2.5e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
 |||||
Db 815 TCCTTGACGTTACTGACGTT 834

RESULT 6

CNS0421L 994 bp DNA GSS 18-MAY-2000
LOCUS
DEFINITION Tetraodon nigroviridis genome survey sequence T7 end of clone
 077D02 of library G from Tetraodon nigroviridis, genomic survey
 sequence.

ACCESSION AL271542 GI:7993521
VERSION
KEYWORDS GSS: genome survey sequence.
SOURCE Tetraodon nigroviridis.
ORGANISM Tetraodon nigroviridis
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
 Eurypterygii; Ctenosquamata; Acanthomorpha; Euacanthomorpha;
 Holacanthopterygii; Acanthopterygii; Percomorpha;
 Tetraodontiformes; Tetraodontidae; Tetraodon.

REFERENCE 1 (bases 1 to 994)
AUTHORS Roest-Crollius,H., Jalllon,O., Dasilva,C., Fizames,C., Fisher,C.,
 Bouneau,L., Billault,A., Quetlier,F., Saurin,W., Bernot,A. and
 Weissenbach,J.
TITLE Characterization and repeat analysis of the compact genome of the
 freshwater pufferfish Tetraodon nigroviridis

JOURNAL Unpublished
REFERENCE 2 (bases 1 to 994)
AUTHORS Roest-Crollius,H., Jalllon,O., Dasilva,C., Bouneau,L., Fisher,C.,
 Bernot,A., Fizames,C., Wincker,P., Brotlier,P., Quetlier,F.,
 Saurin,W. and Weissenbach,J.
TITLE Human gene number estimate provided by genome wide analysis using
 Tetraodon nigroviridis DNA sequence

JOURNAL Unpublished
REFERENCE 3 (bases 1 to 994)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (12-APR-2000) to the EMBL/GenBank/DBJ databases
COMMENT This sequence is a single read and was generated as part of a large
 scale clone-end sequencing project of the Tetraodon nigroviridis
 genome. For more information, please take a look at
 http://www.genoscope.cns.fr/Tetraodon.

FEATURES
source
1..994
/organism="Tetraodon nigroviridis"
/db_xref="taxon:99883"
/clone="077D02"
/clone_1lb="G"
/note="Genoscope sequence ID : C0BG077DB01LP1-end : T7"

BASE COUNT 309 a 199 c 226 g 256 t 4 others
ORIGIN

Query Match 84.0%; Score 16.8; DB 114; length 994;
Best Local Similarity 90.0%; Pred. No. 2.5e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
 |||||
Db 206 TCCATGACGTTCCAGCGTT 225

RESULT 7
CNS01XP2 824 bp DNA GSS 12-MAY-2000
LOCUS
DEFINITION Tetraodon nigroviridis genome survey sequence PUC-Or1 end of clone
 202P06 of library G from Tetraodon nigroviridis, genomic survey
 sequence.

ACCESSION AL172016 GI:7810073
VERSION
KEYWORDS GSS: genome survey sequence.
SOURCE Tetraodon nigroviridis.
ORGANISM Tetraodon nigroviridis
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
 Eurypterygii; Ctenosquamata; Acanthomorpha; Euacanthomorpha;
 Holacanthopterygii; Acanthopterygii; Percomorpha;
 Tetraodontiformes; Tetraodontidae; Tetraodon.

REFERENCE 1 (bases 1 to 824)
AUTHORS Roest-Crollius,H., Jaillon,O., Dasilva,C., Fizames,C., Fisher,C.,
Bouneau,L., Billault,A., Quetier,F., Saurin,W., Bernot,A. and
Weissenbach,J.
TITLE Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 824)
AUTHORS Roest-Crollius,H., Jaillon,O., Dasilva,C., Bouneau,L., Fisher,C.,
Bernot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,
Saurin,W. and Weissenbach,J.
TITLE Human gene number estimate provided by genome wide analysis using
Tetraodon nigroviridis DNA sequence
JOURNAL Unpublished
REFERENCE 3 (bases 1 to 824)
AUTHORS
TITLE Direct Submission
JOURNAL Submitted (12-APR-2000) to the EMBL/GenBank/DBJ databases
COMMENT This sequence is a single read and was generated as part of a large
scale clone-end sequencing project of the Tetraodon nigroviridis
genome. For more information, please take a look at
http://www.genoscope.cns.fr/Tetraodon.
location/Qualifiers
1. 824
/organism="Tetraodon nigroviridis"
/db_xref="taxon:99883"
/clone="202p06"
/clone_lib="G"
/note="Genoscope sequence ID : C0AG202DH03SP1-end :
PUC-ori"
source

BASE COUNT 167 a 236 c 287 g 104 t 30 others
ORIGIN

Query Match 82.0%; Score 16.4; DB 113; Length 824;
Best Local Similarity 94.4%; Pred. No. 3.9e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacg 18
||||| |||||||||
Db 232 TCCATCACGTTCTCTGACG 249

RESULT 8
CNS05NA3/c 1015 bp DNA GSS 26-MAY-2000
LOCUS Tetraodon nigroviridis genome survey sequence SP6 end of clone
DEFINITION 035P01 of library B from Tetraodon nigroviridis, genomic survey
sequence.
ACCESSION AL345108
VERSION AL345108.1 GI:8238878
KEYWORDS GSS; genome survey sequence.
SOURCE Tetraodon nigroviridis.
ORGANISM Tetraodon nigroviridis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Eurypterygii; Ctenosquamata; Acanthomorpha; Euacanthomorpha;
Holacanthopterygii; Acanthopterygii; Percomorpha;
Tetraodontiformes; Tetraodontidae; Tetraodontidae; Tetraodon.
1 (bases 1 to 1015)
Roest-Crollius,H., Jaillon,O., Dasilva,C., Fizames,C., Fisher,C.,
Bouneau,L., Billault,A., Quetier,F., Saurin,W., Bernot,A. and
Weissenbach,J.
TITLE Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 1015)
AUTHORS Roest-Crollius,H., Jaillon,O., Dasilva,C., Bouneau,L., Fisher,C.,
Bernot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,
Saurin,W. and Weissenbach,J.
TITLE Human gene number estimate provided by genome wide analysis using
Tetraodon nigroviridis DNA sequence
JOURNAL Unpublished

REFERENCE 3 (bases 1 to 1015)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (12-APR-2000) to the EMBL/GenBank/DBJ databases
COMMENT This sequence is a single read and was generated as part of a large
scale clone-end sequencing project of the Tetraodon nigroviridis
genome. For more information, please take a look at
http://www.genoscope.cns.fr/Tetraodon.
location/Qualifiers
1. 1015
/organism="Tetraodon nigroviridis"
/db_xref="taxon:99883"
/clone="035P01"
/clone_lib="B"
/note="Genoscope sequence ID : C0B035CH01B1-end : SP6"
source

BASE COUNT 253 a 268 c 261 g 214 t 19 others
ORIGIN

Query Match 82.0%; Score 16.4; DB 115; Length 1015;
Best Local Similarity 94.4%; Pred. No. 4e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 catgacgttcctgacgtt 20
||||| ||||| |||
Db 234 CATGACGTTCTCGATGTT 217

RESULT 9
AV525865 179 bp mRNA EST 21-JUL-2000
LOCUS AV525865 Arabidopsis thaliana aboveground organs two to six-week
DEFINITION old Arabidopsis thaliana cDNA clone APD31f02R 5', mRNA sequence.
ACCESSION AV525865
VERSION AV525865.1 GI:8685393
KEYWORDS EST.
SOURCE thale cress.
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Embryophyta; Tracheophyta; Spermatophyta;
Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids II;
Brassicales; Brassicaceae; Arabidopsis.
1 (bases 1 to 179)
Asamizu,E., Nakamura,Y., Sato,S. and Tabata,S.
A large scale analysis of cDNA in Arabidopsis thaliana: Generation
of 12,028 non-redundant expressed sequence tags from normalized and
size-selected cDNA libraries
DNA Res. 7, 175-180 (2000)
Contact: Erika Asamizu
The First Laboratory for Plant Gene Research
Kazusa DNA Research Institute
Yana 1532-3, Kisarazu, Chiba 292-0812, Japan
Email: asamizu@kazusa.or.jp, URL: http://www.kazusa.or.jp/en/plant/.
location/Qualifiers
1. 179
/organism="Arabidopsis thaliana"
/strain="Columbia"
/db_xref="taxon:3702"
/clone="APD31f02R"
/clone_lib="Arabidopsis thaliana aboveground organs two to
six-week old"
/tissue_type="aboveground organs"
/dev_stage="two to six-week old"
/note="Vector: pBluescriptII SK-; Site_1: EcoRI; Site_2:
XhoI"
source

BASE COUNT 47 a 35 c 48 g 49 t
ORIGIN

Query Match 80.0%; Score 16; DB 19; Length 179;
Best Local Similarity 100.0%; Pred. No. 5.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 tgacgttcctgacgtt 20

Carninci, P. and Hayashizaki, Y.
High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
19-44 (1999)
Please visit our web site (<http://genome.rtc.riken.go.jp>) for
further details.

FEATURES
Location/Qualifiers
1. .262

/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone_lib="C330009E15"
/clone_lib="RIKEN full-length enriched, ES cells"
/cell_type="ES cells"
/lab_host="SOLR"

/note="Site_1: XhoI; Site_2: BamHI; cDNA library was
prepared and sequenced in Mouse Genome Encyclopedia
Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in
RIKEN. Division of Experimental Animal Research in Riken
contributed to prepare mouse tissues. 1st strand cDNA was
primed with a primer [5',
GAGAGAGAGAGATCCAGACCTCTTTTCTTTTCTTTTCTTTVN 3'], cDNA was
prepared by using trehalose thermo-activated reverse
transcriptase and subsequently enriched for full-length by
cap-trapper. cDNA went through one round of normalization
to Rot = 5.0 and subtraction to Rot = 25.0. Second strand
cDNA was prepared with the primer adapter of sequence [5'
GAGAGAGAGATTCGAGTTAATTAAATTAAATCCTCCCCCCCC 3']."

BASE COUNT 53 a 63 c 42 g 103 t 1 others
ORIGIN

Query Match 79.0%; Score 15.8; DB 30; Length 262;
Best Local Similarity 89.5%; Pred. No. 6.9e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 tccatgacgttcctgacgt 19
||||||| |||||
Db 49 TCCATGACACTCCTGACGT 67

RESULT 12
AI077366/c 392 bp mRNA EST 29-SEP-1998
LOCUS
DEFINITION
oy87e11.x1 Soares_fetal_liver_spleen_1NFLS_S1 Homo sapiens cDNA
clone IMAGE:1672844 3', mRNA sequence.

ACCESSION
AI077366
VERSION
AI077366.1 GI:3411774
KEYWORDS
EST.
SOURCE
human.
ORGANISM
Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.

REFERENCE
AUTHORS
TITLE
NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index

JOURNAL
COMMENT
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550

Email: Robert_Strausberg@nih.gov
This clone is available royalty-free through LNL ; contact the
IMAGE Consortium (info@image.lnl.gov) for further information.
Insert length: 904 Std Error: 0.00
Seq primer: -40m13 fwd. ET from Amersham
High quality sequence stop: 294.

FEATURES
Location/Qualifiers
1. .392

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1672844"
/clone_lib="Soares_fetal_liver_spleen_1NFLS_S1"
/sex="male"

/dev_stage="20 week-post conception fetus"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: Liver and Spleen; Vector: pT7T3D (Pharmacia)
with a modified polylinker; Site_1: Pac I; Site_2: Eco RI;
This is a subtracted version of the original Soares fetal
liver spleen 1NFLS library. 1st strand cDNA was primed
with a Pac I - oligo(dT) primer [5',
AACTGAGAGAAATTAATAAGATCTTTTCTTTTCTTTTCTTTT 3'],
double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Pac I and cloned into the Pac I
and Eco RI sites of the modified pT7T3 vector. Library
went through one round of normalization. Library
constructed by Bento Soares and M.Fatima Bonaldo."

BASE COUNT 100 a 86 c 88 g 118 t
ORIGIN

Query Match 79.0%; Score 15.8; DB 8; Length 392;
Best Local Similarity 89.5%; Pred. No. 7.2e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 2 ccatgacgttcctgacgt 20
||||||| |||||
Db 202 CCATGACGTTCCTGAGCGT 184

RESULT 13
AI086210/c 408 bp mRNA EST 28-AUG-1998
LOCUS
DEFINITION
ow90d05.s1 Soares_fetal_liver_spleen_1NFLS_S1 Homo sapiens cDNA
clone IMAGE:1654089 3', mRNA sequence.

ACCESSION
AI086210
VERSION
AI086210.1 GI:3424633
KEYWORDS
EST.
SOURCE
human.
ORGANISM
Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.

REFERENCE
AUTHORS
TITLE
NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index

JOURNAL
COMMENT
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550

Email: Robert_Strausberg@nih.gov
This clone is available royalty-free through LNL ; contact the
IMAGE Consortium (info@image.lnl.gov) for further information.
Insert length: 720 Std Error: 0.00
Seq primer: -40m13 fwd. ET from Amersham
High quality sequence stop: 346.

FEATURES
Location/Qualifiers
1. .408

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1654089"
/clone_lib="Soares_fetal_liver_spleen_1NFLS_S1"
/sex="male"

/dev_stage="20 week-post conception fetus"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: Liver and Spleen; Vector: pT7T3D (Pharmacia)
with a modified polylinker; Site_1: Pac I; Site_2: Eco RI;
This is a subtracted version of the original Soares fetal
liver spleen 1NFLS library. 1st strand cDNA was primed
with a Pac I - oligo(dT) primer [5',
AACTGAGAGAAATTAATAAGATCTTTTCTTTTCTTTTCTTTT 3'],
double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Pac I and cloned into the Pac I
and Eco RI sites of the modified pT7T3 vector. Library
went through one round of normalization. Library
constructed by Bento Soares and M.Fatima Bonaldo."

BASE COUNT 105 a 91 c 94 g 118 t
ORIGIN

Query Match	79.04;	Score 15.8;	DB 8;	Length 408;
Best Local Similarity	89.58;	Pred. No. 7.3e+02;		
Matches 17; Conservative	0;	Mismatches 2;	Indels 0;	Gaps 0;

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QY      2 ccatgacgttcctgacgtt 20
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Db     197 ccATGACGTTCTTGAAGCT 179
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RESULT 14
AW063142/C

LOCUS	AW063142	490 bp	mRNA	EST	01-JUL-2000
DEFINITION	TN0287 KRIBB Human TN Intrathymic T-cell cDNA library Homo sapiens cDNA 3', mRNA sequence.				

ACCESSION	AW063142	
VERSION	AW063142.1	GI:8887191
KEYWORDS	EST.	
SOURCE	human.	

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

AUTHORS

TITLE Expressed sequence tags from three developmental stages of human

JOURNAL

COMMENT

Korea Research Institute of Bioscience and Biotechnology
Oun-dong 52, Yu Sung-Gu, Daejeon 305-333, Republic of Korea
Tel: 82-42-860-4473
Fax: 82-42-860-4479
Email: goshh@mail.kribb.re.kr
Seq primer: T7
High quality sequence stop: 490
POLYA=NO,

High quality sequence stop: 490
POLYA=NO,

FEATURES	Location/Qualifiers
source	1. .490

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/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="KRIBB Human TN Intratymic T-cell CDNA
library"

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/tissue_type="Thymus"
/cell_type="Intrathymic T-cell"
/dev_stage="CD3-4-8- triple negative stage"
/notes="Vector: pGEM-T; cDNA was made from total
cytoplasmic RNA of sorted human intrathymic CD3-4-8-
T-cell, adaptor ligated, amplified with PCR, and cloned
into pGEM-T vector."

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BASE COUNT	103 a	142 c	161 g	83 t	1 others
ORIGIN					

Query Match	79.0%;	Score 15.8;	DB 19;	Length 490;
Best Local Similarity	89.5%;	Pred. NO. 7.4e+02;		
Matches 17;	Conservative	0;	Mismatches 2;	Indels 0;
				Gaps 0;

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QY      2  ccattgacgttcctcgcgctt  20
          || |||||
Db      299 ccgagacgttcctgcagctt  281

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RESULT 1.5

AQ478992

LOCUS	509 bp	DNA	GSS	23-APR-1999
DEFINITION	RPCT-11-254K21.TV RPCT-11 Homo sapiens genomic clone RPCT-11-254K21			

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ACCESSION	AQ478992
VERSION	AQ478992.1
	GI:4661111

KEYWORDS

SOURCE	human.
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ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 509)

REFERENCE	1 (bases 1 to 509)
AUTHORS	Zhao, S., Adams, M.D.

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TITLE

JOURNAL Unpublished (1997)
COMMENT Contact: Shaying Zhao, William Nierman, Mark Adams

The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850

Fax: 301 838 0208

Email: hbeetlgr.org

Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong.

(pieter@edjong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering>) or from Research Genet cs (info@resgen.com). BAC end search page: <http://www.bacpac.org>

Seq primer: r7
Class: BAC ends.

FEATURES	Location/Qualifiers
source	1. .509

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/clone_1fb="RPC1-11"
/sex="Male"
/cell_type="Lymphocytes"
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Best Local Similarity	89.58;	Pred. No. 7.4e+02;		
Matches 17; Conservative	0;	Mismatches 2;	Indels 0;	Gaps 0;

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Db     464 CCATGCCGTTCTGACCTT 482
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Search completed: December 4, 2000, 21:06:41
Job time: 19241 sec

Gencore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 4, 2000, 21:08:04 ; Search time 75.06 Seconds
(without alignments)
40.299 Million cell updates/sec

Title: US-09-369-941-2

Perfect score: 20

Sequence: 1 tccatgacgttcctgacgtt 20

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0
Searched: 262060 seqs, 75620727 residues

Total number of hits satisfying chosen parameters: 524120

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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7: /cgn2_6/ptodata/2/ina/backfiles1.seq:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	20	100.0	20	4	US-09-133-774-12	Sequence 12, Appl
2	20	100.0	20	5	US-09-303-862-12	Sequence 12, Appl
3	16.8	84.0	20	4	US-09-133-774-11	Sequence 11, Appl
4	16.8	84.0	20	5	US-08-386-063-25	Sequence 25, Appl
5	16.8	84.0	20	5	US-09-303-862-11	Sequence 11, Appl
6	16.8	84.0	2470	1	US-07-745-206A-14	Sequence 14, Appl
7	16.8	84.0	2470	3	US-08-311-363-14	Sequence 14, Appl
8	16.8	84.0	5467	1	US-07-745-206A-12	Sequence 12, Appl
9	16.8	84.0	5467	3	US-08-311-363-12	Sequence 12, Appl
10	16.8	84.0	7175	2	US-08-455-543A-8	Sequence 8, Appl
11	16.8	84.0	7175	3	US-08-193-078B-8	Sequence 8, Appl
12	16.8	84.0	7175	3	US-08-223-305C-8	Sequence 8, Appl
13	16.8	84.0	7175	3	US-08-149-097D-8	Sequence 8, Appl
14	16.8	84.0	7175	5	US-08-949-386-8	Sequence 8, Appl
15	16.8	84.0	7175	5	US-08-450-562-8	Sequence 8, Appl
16	16.8	84.0	7266	5	US-08-713-118-1	Sequence 1, Appl
17	16.8	84.0	7362	2	US-08-455-543A-7	Sequence 7, Appl
18	16.8	84.0	7362	3	US-08-193-078B-7	Sequence 7, Appl
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23	15.8	79.0	8931	5	US-09-028-934-28	Sequence 28, Appl
24	15.2	76.0	1799	3	US-08-560-398-3	Sequence 3, Appl
25	14.8	74.0	840	1	US-07-906-983-1	Sequence 1, Appl
26	14.4	72.0	700	5	US-08-764-563-2	Sequence 2, Appl

27	14.2	71.0	29	2	US-08-484-557C-8	Sequence 8, Appl
28	14.2	71.0	29	2	US-08-487-426B-8	Sequence 8, Appl
29	14.2	71.0	29	3	US-08-487-720A-8	Sequence 8, Appl
30	14.2	71.0	76	6	PCT-US96-09451-8	Sequence 8, Appl
31	14.2	71.0	77	5	US-08-945-734-8	Sequence 8, Appl
32	14.2	71.0	355	7	5244792-11	Patent No. 5244792
33	14.2	71.0	523	3	US-08-628-413-1	Sequence 1, Appl
34	14.2	71.0	1949	2	US-08-760-335A-1	Sequence 1, Appl
35	14.2	71.0	1979	2	US-08-392-828C-3	Sequence 3, Appl
36	14.2	71.0	1979	5	US-09-330-945-3	Sequence 3, Appl
37	14.2	71.0	2019	3	US-08-455-073A-5	Sequence 5, Appl
38	14.2	71.0	8140	1	US-08-297-294A-1	Sequence 1, Appl
39	14.2	71.0	50341	2	US-08-247-901C-1	Sequence 1, Appl
40	14.2	71.0	50341	4	US-09-075-904-1	Sequence 17, Appl
41	13.8	69.0	15239	2	US-08-390-878-17	Sequence 7, Appl
42	13.6	68.0	20	1	US-08-436-714-7	Sequence 7, Appl
43	13.6	68.0	20	1	US-08-442-705-7	Sequence 7, Appl
44	13.6	68.0	20	2	US-08-332-829-7	Sequence 7, Appl
45	13.6	68.0	20	5	US-08-386-063-21	Sequence 21, Appl

ALIGNMENTS

RESULT 1
US-09-133-774-12
; Sequence 12, Application US/09133774B
; Patent No. 5962636
; GENERAL INFORMATION:
; APPLICANT: Bachmaier, Kurt
; APPLICANT: Hessel, Andrew J.
; APPLICANT: Neu M.D., Nikolaus
; APPLICANT: Penninger, Josef M.
; TITLE OF INVENTION: No. 5962636el Peptides Capable of Modulating Inflammatory Hear
; TITLE OF INVENTION: Disease
; FILE REFERENCE: A-536
; CURRENT APPLICATION NUMBER: US/09/133,774B
; CURRENT FILING DATE: 1998-08-12
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 12
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Chlamydia trachomatis
; FEATURE:
; OTHER INFORMATION: An oligonucleotide derived from the DNA encoding a
; OTHER INFORMATION: 60 kda cysteine rich outer membrane protein from
; OTHER INFORMATION: Chlamydia trachomatis.
US-09-133-774-12

Query Match 100.0%; Score 20; DB 4; Length 20;
Best local Similarity 100.0%; Pred. No. 0.089;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 1 tccatgacgttcctgacgtt 20

RESULT 2
US-09-303-862-12
; Sequence 12, Application US/09303862
; Patent No. 6034230
; GENERAL INFORMATION:
; APPLICANT: Bachmaier, Kurt
; APPLICANT: Hessel, Andrew J.
; APPLICANT: Neu M.D., Nikolaus
; APPLICANT: Penninger, Josef M.
; TITLE OF INVENTION: No. 6034230el Peptides Capable of Modulating Inflammatory Hear
; TITLE OF INVENTION: Disease
; FILE REFERENCE: A-536
; CURRENT APPLICATION NUMBER: US/09/303,862

Db 1 tccatgacgttcctgatgt 20

RESULT 6

US-07-745-206A-14/c

; Sequence 14, Application US/07745206A

; Patent No. 5429921

; GENERAL INFORMATION:

; APPLICANT: Harpold, Michael

; APPLICANT: Ellis, Steven

; APPLICANT: Williams, Mark

; APPLICANT: McCue, Ann

; APPLICANT: Feldman, Daniel

; TITLE OF INVENTION: Human Calcium Channel Compositions and

; TITLE OF INVENTION: Methods

; NUMBER OF SEQUENCES: 32

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Fitch, Even, Tabin & Flannery

; STREET: 135 S. LaSalle

; CITY: Chicago

; STATE: Illinois

; COUNTRY: U.S.A.

; ZIP: 60603

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: PatentIn Release #1.0, Version #1.25

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/07/745,206A

; FILING DATE: 19910815

; CLASSIFICATION: 435

; ATTORNEY/AGENT INFORMATION:

; NAME: Feder, Scott B

; REFERENCE/DOCKET NUMBER: 51504

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: 312-372-7842

; INFORMATION FOR SEQ ID NO: 14:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 2470 base pairs

; TYPE: NUCLEIC ACID

; STRANDEDNESS: unknown

; TOPOLOGY: unknown

; MOLECULE TYPE: DNA (genomic)

; FEATURE:

; NAME/KEY: CDS

; LOCATION: 1..2469

US-07-745-206A-14

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Best Local Similarity 90.0%; Pred. No. 6.6;

Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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Db 509 TCCATGACGTTCCAGCCGTT 490

RESULT 7

US-08-311-363-14/c

; Sequence 14, Application US/08311363

; Patent No. 5876958

; GENERAL INFORMATION:

; APPLICANT: Harpold, Michael

; APPLICANT: Ellis, Steven

; APPLICANT: Williams, Mark

; APPLICANT: Feldman, Daniel

; APPLICANT: McCue, Ann

; APPLICANT: Brenner, Robert

; TITLE OF INVENTION: Human Calcium Channel Compositions and

; TITLE OF INVENTION: Methods

; NUMBER OF SEQUENCES: 32

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Brown, Martin, Haller & McClain

; STREET: 1660 Union Street

; CITY: San Diego

; STATE: California

; COUNTRY: USA

; ZIP: 92101-2926

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: PatentIn Release #1.0, Version #1.25

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/311,363

; FILING DATE:

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: US 07/745,206

; FILING DATE: 15-AUG-1991

; ATTORNEY/AGENT INFORMATION:

; NAME: Seidman, Stephanie L.

; REGISTRATION NUMBER: 33,779

; REFERENCE/DOCKET NUMBER: 6362-51506

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (619)238-0999

; TELEFAX: (619)238-0062

; INFORMATION FOR SEQ ID NO: 14:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 2470 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: unknown

; TOPOLOGY: unknown

; MOLECULE TYPE: DNA (genomic)

; FEATURE:

; NAME/KEY: CDS

; LOCATION: 1..2469

US-08-311-363-14

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Best Local Similarity 90.0%; Pred. No. 6.6;

Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgatgt 20

|||||

Db 509 TCCATGACGTTCCAGCCGTT 490

RESULT 8

US-07-745-206A-12/c

; Sequence 12, Application US/07745206A

; Patent No. 5429921

; GENERAL INFORMATION:

; APPLICANT: Harpold, Michael

; APPLICANT: Ellis, Steven

; APPLICANT: Williams, Mark

; APPLICANT: McCue, Ann

; APPLICANT: Feldman, Daniel

; TITLE OF INVENTION: Human Calcium Channel Compositions and

; TITLE OF INVENTION: Methods

; NUMBER OF SEQUENCES: 32

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Fitch, Even, Tabin & Flannery

; STREET: 135 S. LaSalle

; CITY: Chicago

; STATE: Illinois

; COUNTRY: U.S.A.

; ZIP: 60603

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: PatentIn Release #1.0, Version #1.25

; CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/07/745,206A
FILING DATE: 19910815
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Feder, Scott B
REFERENCE/DOCKET NUMBER: 51504
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312-372-7842
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 5467 base pairs
TYPE: NUCLEIC ACID
STRANDEDNESS: unknown
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: join(144..3164, 3168..3245, 3249..3386, 3390
LOCATION: ..3392, 3396..3488, 3495..3539, 3543..3581, 3585
LOCATION: ..3587, 3591..3626, 3630..3689, 3693..3737, 3744
LOCATION: ..3746, 3750..4823, 4827..4841, 4845..5006, 5010
LOCATION: ..5096, 5100..5306, 5310..5366, 5370..5465)
US-07-745-206A-12

Query Match 84.0%; Score 16.8; DB 1; Length 5467;
Best Local Similarity 90.0%; Pred. No. 7.3;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
|||||
Db 652 tccatgacgttccagccgtt 633

RESULT 9
US-08-311-363-12/c
Sequence 12, Application US/08311363
Patent No. 5876958
GENERAL INFORMATION:
APPLICANT: Harpold, Michael
APPLICANT: Ellis, Steven
APPLICANT: Williams, Mark
APPLICANT: Feldman, Daniel
APPLICANT: McCue, Ann
APPLICANT: Brenner, Robert
TITLE OF INVENTION: Human Calcium Channel Compositions and
TITLE OF INVENTION: Methods
NUMBER OF SEQUENCES: 32
CORRESPONDENCE ADDRESS:
ADDRESSEE: Brown, Martin, Haller & McClain
STREET: 1660 Union Street
CITY: San Diego
STATE: California
COUNTRY: USA
ZIP: 92101-2926
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/311,363
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/745,206
FILING DATE: 15-AUG-1991
ATTORNEY/AGENT INFORMATION:
NAME: Seidman, Stephanie L.
REGISTRATION NUMBER: 33,779
REFERENCE/DOCKET NUMBER: 6362-51506
TELECOMMUNICATION INFORMATION:
TELEPHONE: (619)238-0999
TELEFAX: (619)238-0062

INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 5467 base pairs
TYPE: nucleic acid
STRANDEDNESS: unknown
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: join(144..3164, 3168..3245, 3249..3386, 3390
LOCATION: ..3392, 3396..3488, 3495..3539, 3543..3581, 3585
LOCATION: ..3587, 3591..3626, 3630..3689, 3693..3737, 3744
LOCATION: ..3746, 3750..4823, 4827..4841, 4845..5006, 5010
LOCATION: ..5096, 5100..5306, 5310..5366, 5370..5465)
US-08-311-363-12

Query Match 84.0%; Score 16.8; DB 3; Length 5467;
Best Local Similarity 90.0%; Pred. No. 7.3;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
|||||
Db 652 tccatgacgttccagccgtt 633

RESULT 10
US-08-455-543A-8/c
Sequence 8, Application US/08455543A
Patent No. 5792846
GENERAL INFORMATION:
APPLICANT: Harpold, Michael
APPLICANT: Ellis, Steven
APPLICANT: Williams, Mark
APPLICANT: Feldman, Daniel
APPLICANT: McCue, Ann
APPLICANT: Brenner, Robert
TITLE OF INVENTION: HUMAN CALCIUM CHANNEL COMPOSITIONS AND
TITLE OF INVENTION: METHODS
NUMBER OF SEQUENCES: 57
CORRESPONDENCE ADDRESS:
ADDRESSEE: Brown, Martin, Haller & McClain
STREET: 1660 Union Street
CITY: San Diego
STATE: California
COUNTRY: USA
ZIP: 92101-2926
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/455,543A
FILING DATE: May 31, 1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/223,305
FILING DATE: April 4, 1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/868,354
FILING DATE: April 10, 1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/745,206
FILING DATE: 15-AUG-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/620,250
FILING DATE: 30-NOV-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/482,384
FILING DATE: 20-FEB-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/603,751
FILING DATE: 04-APR-1989

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; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: WO PCT/US89/01408
; FILING DATE: 04-APR-1989
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/176,899
; FILING DATE: 04-APR-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Seidman, Stephanie L.
; REGISTRATION NUMBER: 33,779
; REFERENCE/DOCKET NUMBER: 6362-52517
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619)238-0999
; TELEFAX: (619)238-0062
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7175 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 144..6857
; FEATURE:
; NAME/KEY: 5'UTR
; LOCATION: 1..143
; FEATURE:
; NAME/KEY: 3'UTR
; LOCATION: 6855..7175
; US-08-455-543A-8
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Query Match      84.0%; Score 16.8; DB 2; Length 7175;
Best Local Similarity 90.0%; Pred. No. 7.6;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
    |||||1111111111111111
Db 652 TCCATGACGTTCACGCCGTT 633
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RESULT 11
US-08-193-078B-8/c
; Sequence 8, Application US/08193078B
; Patent No. 5846757
; GENERAL INFORMATION:
; APPLICANT: Harpold, Michael
; APPLICANT: Ellis, Steven
; APPLICANT: Williams, Mark
; APPLICANT: Feldman, Daniel
; APPLICANT: McCue, Ann
; APPLICANT: Brenner, Robert
; TITLE OF INVENTION: HUMAN CALCIUM CHANNEL COMPOSITIONS AND
; TITLE OF INVENTION: METHODS
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BROWN, MARTIN, HALLER & McCLAIN
; STREET: 1660 UNION STREET
; CITY: SAN DIEGO
; STATE: CA
; COUNTRY: USA
; ZIP: 92101
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/193,078B
; FILING DATE: 07-FEB-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/868,354
```

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; FILING DATE: 10-APR-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/745,206
; FILING DATE: 15-AUG-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Seidman, Stephanie L.
; REGISTRATION NUMBER: 33,779
; REFERENCE/DOCKET NUMBER: 6362-53607
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 619-238-0999
; TELEFAX: 619-238-0062
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7175 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 144..6857
; FEATURE:
; NAME/KEY: 5'UTR
; LOCATION: 1..143
; FEATURE:
; NAME/KEY: 3'UTR
; LOCATION: 6855..7175
; US-08-193-078B-8
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Query Match      84.0%; Score 16.8; DB 3; Length 7175;
Best Local Similarity 90.0%; Pred. No. 7.6;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
    |||||1111111111111111
Db 652 TCCATGACGTTCACGCCGTT 633
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RESULT 12
US-08-223-305C-8/c
; Sequence 8, Application US/08223305C
; Patent No. 5851824
; GENERAL INFORMATION:
; APPLICANT: Harpold, Michael
; APPLICANT: Ellis, Steven
; APPLICANT: Williams, Mark
; APPLICANT: Feldman, Daniel
; APPLICANT: McCue, Ann
; APPLICANT: Brenner, Robert
; TITLE OF INVENTION: HUMAN CALCIUM CHANNEL COMPOSITIONS AND
; TITLE OF INVENTION: METHODS
; NUMBER OF SEQUENCES: 57
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Brown, Martin, Haller & McClain
; STREET: 1660 Union Street
; CITY: San Diego
; STATE: California
; COUNTRY: USA
; ZIP: 92101-2926
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq Version 1.5
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/223,305C
; FILING DATE: April 4, 1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/868,354
; FILING DATE: April 10, 1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/745,206
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; FILING DATE: 15-AUG-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/620,250
; FILING DATE: 30-NOV-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/482,384
; FILING DATE: 20-FEB-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/603,751
; FILING DATE: 04-APR-1989
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: WO PCT/US89/01408
; FILING DATE: 04-APR-1989
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/176,899
; FILING DATE: 04-APR-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Seidman, Stephanie L.
; REGISTRATION NUMBER: 33,779
; REFERENCE/DOCKET NUMBER: 52516 (P519739)
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619)238-0999
; TELEFAX: (619)238-0062
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7175 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 144..6857
; FEATURE:
; NAME/KEY: 5'UTR
; LOCATION: 1..143
; FEATURE:
; NAME/KEY: 3'UTR
; LOCATION: 6855..7175
; US-08-223-305C-8

Query Match      84.0%; Score 16.8; DB 3; Length 7175;
Best Local Similarity 90.0%; Pred. No. 7.6;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
    |||||
Db 652 tccatgacgttccagccgtt 633

RESULT 13
US-08-149-097D-8/c
; Sequence 8, Application US/08149097D
; Patent No. 5874236
; GENERAL INFORMATION:
; APPLICANT: Harpold, Michael
; APPLICANT: Ellis, Steven
; APPLICANT: Williams, Mark
; APPLICANT: Feldman, Daniel
; APPLICANT: McCue, Ann
; APPLICANT: Brenner, Robert
; TITLE OF INVENTION: HUMAN CALCIUM CHANNEL COMPOSITIONS AND
; NUMBER OF SEQUENCES: 40
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Brown, Martin, Haller & McClain
; STREET: 1660 Union Street
; CITY: San Diego
; STATE: California
; COUNTRY: USA
; ZIP: 92101-2926
; COMPUTER READABLE FORM:
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; MEDIUM TYPE: floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/149,097D
; FILING DATE: 05-NOV-1993
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/105,536
; FILING DATE: 11-AUG-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: WO PCT/US92/06903
; FILING DATE: 14-AUG-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/914,231
; FILING DATE: 13-JUL-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/868,354
; FILING DATE: 10-APR-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/745,206
; FILING DATE: 15-AUG-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/620,250
; FILING DATE: 30-NOV-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/482,384
; FILING DATE: 20-FEB-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/603,751
; FILING DATE: 04-APR-1989
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: WO PCT/US89/01408
; FILING DATE: 04-APR-1989
; APPLICATION NUMBER: US 07/176,899
; FILING DATE: 04-APR-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Seidman, Stephanie L.
; REGISTRATION NUMBER: 33,779
; REFERENCE/DOCKET NUMBER: 6362-55038
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 238-0999
; TELEFAX: (619) 238-0062
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7175 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 144..6857
; FEATURE:
; NAME/KEY: 5'UTR
; LOCATION: 1..143
; FEATURE:
; NAME/KEY: 3'UTR
; LOCATION: 6855..7175
; US-08-149-097D-8

Query Match      84.0%; Score 16.8; DB 3; Length 7175;
Best Local Similarity 90.0%; Pred. No. 7.6;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
    |||||
Db 652 tccatgacgttccagccgtt 633
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RESULT 14
US-08-949-386-8/c
; Sequence 8, Application US/08949386
; Patent No. 6090623
; GENERAL INFORMATION:
; APPLICANT: Harpold, Michael
; APPLICANT: Ellis, Steven
; APPLICANT: Williams, Mark
; APPLICANT: McCue, Ann
; APPLICANT: Gillespie, Alison
; TITLE OF INVENTION: HUMAN CALCIUM CHANNEL COMPOSITIONS AND
; TITLE OF INVENTION: METHODS
; NUMBER OF SEQUENCES: 38
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Brown, Martin, Haller & McClain
; STREET: 1660 Union Street
; CITY: San Diego
; STATE: California
; COUNTRY: US
; ZIP: 92101
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/949,386
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/290,012
; FILING DATE: 11-AUG-1994
; APPLICATION NUMBER: 08/149,097
; FILING DATE: 5-NOV-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/105,536
; FILING DATE: 11-AUG-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Seidman, Stephanie L.
; REGISTRATION NUMBER: 33,779
; REFERENCE/DOCKET NUMBER: 519808
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 238-0999
; TELEFAX: (619) 238-0062
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7175 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 144..6857
; FEATURE:
; NAME/KEY: 5'UTR
; LOCATION: 1..143
; FEATURE:
; NAME/KEY: 3'UTR
; LOCATION: 6855..7175
; US-08-949-386-8

Query Match 84.0%; Score 16.8; DB 5; Length 7175;
Best Local Similarity 90.0%; Pred. No. 7.6;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
|||||
Db 652 TCCATGACGTTCCAGCCGTT 633

RESULT 15

US-08-450-562-8/c
; Sequence 8, Application US/08450562
; Patent No. 6096514
; GENERAL INFORMATION:
; APPLICANT: Harpold, Michael
; APPLICANT: Ellis, Steven
; APPLICANT: Williams, Mark
; APPLICANT: McCue, Ann
; APPLICANT: Gillespie, Alison
; APPLICANT: Feldman, Daniel
; APPLICANT: Brenner, Robert
; TITLE OF INVENTION: HUMAN CALCIUM CHANNEL COMPOSITIONS AND
; TITLE OF INVENTION: METHODS
; NUMBER OF SEQUENCES: 38
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Brown, Martin, Haller & McClain
; STREET: 1660 Union Street
; CITY: San Diego
; STATE: California
; COUNTRY: US
; ZIP: 92101
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/450,562
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/404,950
; FILING DATE: 13-MAR-1995
; APPLICATION NUMBER: 08/336,257
; FILING DATE: 7-NOV-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/314,083
; FILING DATE: 28-SEPT-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/311,363
; FILING DATE: 23-SEPT-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/290,012
; FILING DATE: 11-AUG-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/223,305
; FILING DATE: 4-APR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/193,078
; FILING DATE: 07-FEB-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/149,097
; FILING DATE: 5-NOV-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/105,536
; FILING DATE: 11-AUG-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/914,231
; FILING DATE: 13-JULY-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/868,354
; FILING DATE: 10-APR-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US92/06903
; FILING DATE: 14-AUG-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/745,206
; FILING DATE: 15-AUG-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/620,250
; FILING DATE: 30-NOV-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/603,751


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1      FILING DATE: 08-NOV-1990
2      PRIOR APPLICATION DATA:
3      APPLICATION NUMBER: 07/482,384
4      FILING DATE: 02-FEB-1990
5      PRIOR APPLICATION DATA:
6      APPLICATION NUMBER: PCT/US89/01408
7      FILING DATE: 04-APR-1989
8      PRIOR APPLICATION DATA:
9      APPLICATION NUMBER: 07/176,899
10     FILING DATE: 04-APR-1988
11     ATTORNEY/AGENT INFORMATION:
12     NAME: Seidman, Stephanie L.
13     REGISTRATION NUMBER: 33,779
14     REFERENCE/DOCKET NUMBER: 6362-519812
15     TELECOMMUNICATION INFORMATION:
16     TELEPHONE: (619) 238-0999
17     TELEFAX: (619) 238-0062
18     INFORMATION FOR SEQ ID NO: 8:
19     SEQUENCE CHARACTERISTICS:
20     LENGTH: 7175 base pairs
21     TYPE: nucleic acid
22     STRANDEDNESS: double
23     TOPOLOGY: linear
24     MOLECULE TYPE: DNA (genomic)
25     FEATURE:
26     NAME/KEY: CDS
27     LOCATION: 144..6857
28     FEATURE:
29     NAME/KEY: 5'UTR
30     LOCATION: 1..143
31     FEATURE:
32     NAME/KEY: 3'UTR
33     LOCATION: 6855..7175
34
35     US-08-450-562-8

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Query Match	84.0%;	Score 16.8;	DB 5;	Length 7175;
Best Local Similarity	90.0%;	Pred. No. 7.6;		
Matches 18;	Conservative 0;	Mismatches 2;	Indels 0;	Gaps 0;

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QY      1 tccatgacgttcctgacgt 20
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Db      652 TCCATGACGTTCCAGCCGTT 633

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Search completed: December 4, 2000, 21:08:06
Job time: 16804 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 4, 2000, 21:09:55 ; Search time 99.31 Seconds
(without alignments)
75.655 Million cell updates/sec

Title: US-09-369-941-2
Perfect score: 20
Sequence: 1 tccatgacgttcctgacgtt 20

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 480022 seqs, 187831343 residues

Total number of hits satisfying chosen parameters: 960044

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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15: /cgn2_2/gcgdata/geneseq/geneseqn/NA1994.DAT:*
16: /cgn2_2/gcgdata/geneseq/geneseqn/NA1995.DAT:*
17: /cgn2_2/gcgdata/geneseq/geneseqn/NA1996.DAT:*
18: /cgn2_2/gcgdata/geneseq/geneseqn/NA1997.DAT:*
19: /cgn2_2/gcgdata/geneseq/geneseqn/NA1998.DAT:*
20: /cgn2_2/gcgdata/geneseq/geneseqn/NA1999.DAT:*
21: /cgn2_2/gcgdata/geneseq/geneseqn/NA2000.DAT:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	20	100.0	20	19	V60950	unmethylated cytos
2	20	100.0	20	19	V47683	unmethylated Cpg d
3	20	100.0	20	19	V27667	Immunostimulatory
4	20	100.0	20	20	Z31943	Cpg adjuvant oligo
5	20	100.0	20	20	Z41946	IL-12 secretion in
6	20	100.0	20	20	Z41949	IL-12 secretion in
7	20	100.0	20	20	Z28191	Chlamydia trachoma
8	20	100.0	20	20	X78802	HPV fusion protein
9	20	100.0	20	20	X88536	Cytosine-guanosine
10	20	100.0	20	20	V74237	Cpg-N motif S-ODN
11	20	100.0	20	20	V74242	Cpg-N motif O-ODN
12	20	100.0	20	20	V74244	Cpg-N motif SOS-OD

13	20	100.0	20	21	Z99004	Cpg motif for immu
14	20	100.0	20	21	Z99174	Inflammatory cardi
15	20	100.0	20	21	Z61010	Nucleotide sequenc
16	20	100.0	20	21	Z61012	Nucleotide sequenc
17	20	100.0	20	21	Z47601	Murine immune syst
18	20	100.0	20	21	Z47885	Immunostimulatory
19	20	100.0	20	21	Z47887	Immunostimulatory
20	20	100.0	20	21	Z48022	Immune remodeling
21	20	100.0	20	21	Z48025	Immune remodeling
22	20	100.0	44	20	V83723	Murine-specific Cp
23	20	100.0	44	20	V83722	Murine-specific Cp
24	20	100.0	44	20	V83726	Cpg-optimised gene
25	17	85.0	17	19	V52557	Unmethylated Cpg d
26	17	85.0	17	19	V27731	Immunostimulatory
27	17	85.0	17	20	Z41916	IL-12 secretion in
28	17	85.0	17	21	Z60984	Nucleotide sequenc
29	17	85.0	17	21	Z47653	Parasitic infectio
30	17	85.0	17	21	Z47859	Immunostimulatory
31	17	85.0	17	21	Z47992	Immune remodeling
32	16.8	84.0	20	18	T88792	Synthetic phosphor
33	16.8	84.0	20	19	V52567	Unmethylated Cpg d
34	16.8	84.0	20	19	V45995	Immune adjuvant Cp
35	16.8	84.0	20	19	V45996	Immune adjuvant Cp
36	16.8	84.0	20	19	V27708	Immunostimulatory
37	16.8	84.0	20	19	V27700	Immunostimulatory
38	16.8	84.0	20	19	V27651	Immunostimulatory
39	16.8	84.0	20	19	V27638	Immunostimulatory
40	16.8	84.0	20	20	Z41879	IL-12 secretion in
41	16.8	84.0	20	20	Z41919	IL-12 secretion in
42	16.8	84.0	20	20	Z41930	IL-12 secretion in
43	16.8	84.0	20	20	Z28190	Chlamydia trachoma
44	16.8	84.0	20	20	V72500	Cpg motif containi
45	16.8	84.0	20	20	V74261	Cpg-N motif oligon

ALIGNMENTS

RESULT	ID	Description
1	V60950	standard; DNA; 20 BP.
XX	XX	
AC	V60950;	
XX	XX	
DT	14-DEC-1998	(first entry)
XX	XX	
DE	Unmethylated cytosine-guanine dinucleotide containing oligonucleotide 1.	
XX	XX	
KW	ss; unmethylated Cpg dinucleotide; immune response; natural killer cell;	
KW	Th2 response; Th1 response; Th1 cytokine; hepatitis B.	
XX	XX	
OS	Synthetic.	
XX	XX	
PN	WO9840100-A1.	
XX	XX	
PD	17-SEP-1998.	
XX	XX	
PF	10-MAR-1998;	98WO-US04703.
XX	XX	
PR	10-MAR-1997;	97US-0040376.
XX	XX	
PA	(OTTA-) OTTAWA CIVIC LOEB RES INST.	
PA	(QIAG-) QIAGEN GMBH.	
PA	(IOWA) UNIV IOWA RES FOUND.	
XX	XX	
PI	Davis HL, Krieg AM, Schorr J;	
XX	XX	
DR	WPI; 1998-520792/44.	
XX	XX	
PT	Use of oligonucleotides containing an unmethylated Cpg dinucleotide	
PT	- useful as, e.g. adjuvant with antigen, or nucleic acid encoding	
PT	antigen for inducing immune response in subject	
XX	XX	

```
PS Claim 14; Page 35; 67pp; English.
XX
CC Oligonucleotides containing at least 1 unmethylated Cpg dinucleotide
CC affect the immune response in a subject by activating natural killer
CC cells or redirecting a subject's immune response from a Th2 to a Th1
CC response by inducing monocytic and other cells to produce Th1 cytokines.
CC These nucleic acids containing at least 1 unmethylated Cpg can be used as
CC an adjuvant, specifically to induce an immune response against an
CC antigenic protein, and are used particularly for virally mediated
CC disorders, e.g. hepatitis B virus infection.
XX
SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match          100.0%; Score 20; DB 19; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
   |||||
DB 1 tccatgacgttcctgacgtt 20

RESULT 2
V47683 V47683 standard; DNA; 20 BP.
XX
AC V47683;
XX
DT 20-NOV-1998 (first entry)
XX
DE Unmethylated Cpg dinucleotide 1826.
XX
KW Unmethylated Cpg dinucleotide; immune response; bacterial meningitis;
KW natural killer cell activation; NK cell; Th2 response; neonatal sepsis;
KW pulmonary disorder; asthma; environmentally induced airway disease;
KW bacterial infection; endotoxaemia; therapy; cystic fibrosis;
KW inflammatory bowel disease; ss.
XX
OS Synthetic.
XX
PN WO9837919-A1.
XX
PD 03-SEP-1998.
XX
PF 25-FEB-1998; 98WO-US03678.
XX
PR 28-FEB-1997; 97US-0039405.
XX
PA (IOWA ) UNIV IOWA RES FOUND.
XX
PI Krieg AM, Schwartz DA;
XX
PI Krieg AM, Schwartz DA;
XX
DR WPI; 1998-480941/41.
XX
PT Use of nucleic acids containing an unmethylated Cpg - for treating a
PT subject having or at risk of having an acute decrement in air flow
PT or inhibiting an inflammatory response
XX
PS Claim 35; Page 27; 65pp; English.
XX
CC This sequence represents an unmethylated Cpg dinucleotide, and can be
CC used in the method of the invention. The method is for treating a subject
CC having, or at risk of having an acute decrement in air flow, comprising
CC administering a nucleic acid sequence containing at least one
CC unmethylated Cpg. The nucleic acids containing an unmethylated Cpg
CC dinucleotide affect an immune response in a subject by activating natural
CC killer cells (NK) or redirecting a subject's immune response from a Th2
CC to a Th1 response by inducing monocytic and other cells to produce Th1
CC cytokines. They can be used to treat pulmonary disorders having an
CC immunologic component, such as asthma or environmentally induced airway
CC disease. They can also be used to treat diseases associated with
CC Gram-positive bacterial infections or endotoxaemia including bacterial
```

```
CC meningitis, neonatal sepsis, cystic fibrosis, inflammatory bowel disease
CC and liver cirrhosis, Gram-negative pneumonia, Gram-negative abdominal
CC abscess, haemorrhagic shock, disseminated intravascular coagulation, or
CC an inflammatory response to lipopolysaccharide.
XX
SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match          100.0%; Score 20; DB 19; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
   |||||
DB 1 tccatgacgttcctgacgtt 20

RESULT 3
V27667 V27667 standard; DNA; 20 BP.
XX
AC V27667;
XX
DT 01-OCT-1998 (first entry)
XX
DE Immunostimulatory oligodeoxyribonucleotide of the invention.
XX
KW Immunostimulatory; oligodeoxyribonucleotide; ODN;
KW unmethylated Cpg dinucleotide; activate; lymphocyte; immune response;
KW Th2; Th1; cytokine; treatment; prevention; asthma; autoimmune disease;
KW desensitisation therapy; artificial adjuvant; antibody generation; ss.
XX
OS Synthetic.
XX
PN WO9818810-A1.
XX
PD 07-MAY-1998.
XX
PF 30-OCT-1997; 97WO-US19791.
XX
PR 30-OCT-1996; 96US-0738652.
XX
PA (IOWA ) UNIV IOWA RES FOUND.
XX
PI Kline JN, Krieg AM;
XX
PI Kline JN, Krieg AM;
XX
DR WPI; 1998-272127/24.
XX
PT New immunostimulatory nucleic acid molecules - which contain at
PT least one unmethylated Cpg dinucleotide, used for treating e.g.
PT tumours, infections or autoimmune disease
XX
PS Claim 35; Page 84; 109pp; English.
XX
CC V27641-751 represent immunostimulatory oligodeoxyribonucleotides (ODNs)
CC of the invention. The ODNs contain at least one unmethylated Cpg
CC dinucleotide, and have the formula:
CC 5' N1X1CGX2N2 3', where at least one nucleotide separates consecutive
CC Cpgs, X1 is adenine, guanine, or thymine, X2 is cytosine or thymine, N is
CC any nucleotide and N1+N2 is 0-26 bases with the provision that N1 and N2
CC does not contain a CCGG tetramer or more than one CCG or CGG trimer OR
CC 5' NX1X2CGX3X4N 3', where at least one nucleotide separates consecutive
CC Cpgs, X1 and X2 are selected from GpT, GpG, GpA, ApT and ApA, X3and X4
CC are selected from TpT or CpT, N is any nucleotide and N1+N2 is 0-26
CC bases with the provision that N1 and N2 does not contain a CCGG tetramer
CC or more than one CCG or CGG trimer.
CC The ODNs activate lymphocytes in a subject and redirect a subject's
CC immune response from a Th2 to a Th1 (e.g. by inducing monocytic cells and
CC other cells to produce Th1 cytokines, including IL-12, IFN-gamma and
CC GM-CSF). The ODNs can be used to treat or prevent an asthmatic disorder,
CC autoimmune diseases, in desensitisation therapy, as an artificial
CC adjuvant during antibody generation in a mammal such as a mouse or a
CC human.
```


XX Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;
SQ

Query Match 100.0%; Score 20; DB 19; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 tccatgacgttcctgacgtt 20
|||||
Db 1 tccatgacgttcctgacgtt 20

RESULT 4

Z31943
ID Z31943 standard; DNA; 20 BP.

AC Z31943;

DT 26-JAN-2000 (first entry)

DE Cpg adjuvant oligo 1001.

KW Cpg adjuvant; vaccine; polyoxyethylene ether; polyoxyethylene ester;
antigen; infection; allergy; cancer; therapy; ss.

OS Synthetic.

PN WO99512549-A1.

PD 21-OCT-1999.

PF 29-MAR-1999; 99WO-EP02278.

PR 09-APR-1998; 98GB-0007805.

PR 25-SEP-1998; 98GB-0020956.

PA (SMIK) SMITHKLINE BEECHAM BIOLOGICALS.

PI Friede M, Hermand P;

DR WPI; 1999-620290/53.

PT Vaccine to protect against infections, allergy and cancer -

PS Claim 16; Page 32; 52pp; English.

CC This sequence represents a Cpg adjuvant that can be used in the vaccine
CC composition of the invention. The vaccine comprises a polyoxyethylene
CC ether or ester (I), not in the form of a vesicle, pharmaceutically
CC acceptable excipient and an antigen (Ag) or antigenic composition. The
CC vaccine can be used to treat or prevent infections (by bacteria, viruses
CC or other parasites), allergy and cancer. (I), which are safe, easy to
CC sterilize and simple to administer, are powerful vaccine adjuvants, able
CC to induce a systemic immune response when administered (non-invasively)
CC to the mucosa. The response is at least as good as that from conventional
CC systemic injection. (I) are effective at low concentration, have low
CC reactogenicity and are well tolerated.

SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match 100.0%; Score 20; DB 20; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 tccatgacgttcctgacgtt 20
|||||
Db 1 tccatgacgttcctgacgtt 20

RESULT 5
Z41946

ID Z41946 standard; DNA; 20 BP.

AC Z41946;

DT 24-JAN-2000 (first entry)

DE IL-12 secretion inducing Cpg oligonucleotide 91.

KW Cpg oligonucleotide; phosphorothioate; interleukin-12; IL-12; secretion;
human PBMC; immune response; cancer; HIV; bacterial disease; asthma;
neoplastic disorder; jaagsiekte; B cell; NK cell; ss; cytokine;
antigen presenting cell; infection; allergic disease.

OS Synthetic.

PN WO9951259-A2.

PD 14-OCT-1999.

PF 02-APR-1999; 99WO-US07335.

PR 03-APR-1998; 98US-0080729.

PA (IOWA) UNIV IOWA RES FOUND.

PI Krieg AM, Weiner G;

DR WPI; 1999-620169/53.

PT Novel synergistic combinations of immunostimulatory oligonucleotides
and immunopotentiating cytokines are useful for stimulating the immune
system -

PS Example 8; Page 88; 91pp; English.

CC Sequences Z41856-Z41949 are phosphorothioate Cpg oligonucleotides which
CC are used in the invention to induce interleukin-12 (IL-12) secretion
CC from human PBMC. The invention comprises stimulating an immune response
CC in a subject comprising administering to a subject exposed to an antigen,
CC an immunopotentiating cytokine and an immunostimulatory Cpg
CC oligonucleotide to induce a synergistic antigen specific immune response.
CC The methods are useful for treating cancer by stimulating an antigen
CC specific immune response against a cancer antigen. The methods can also
CC be used to treat neoplastic disorders in humans, including but not
CC limited to: sarcoma, carcinoma, fibroma, lymphoma, melanoma,
CC neuroblastoma, retinoblastoma, and glioma. The methods are also useful
CC for treating infectious diseases, e.g. viral diseases such as HIV,
CC bacterial diseases, and fungal diseases. The methods may also be used to
CC treat allergic diseases, e.g. asthma. The methods and compositions may
CC also be applied to treat cancer and tumours in non human subjects,
CC e.g. cats and dogs. Neoplasias affecting agricultural livestock may also
CC be treated and include leukaemia, haemangiopericytoma and bovine ocular
CC neoplasia. Chronic, infectious, contagious diseases of sheep and goats
CC caused by the bacterium Corynebacterium pseudotuberculosis, and
CC contagious lung tumour of sheep caused by jaagsiekte may also be treated.
CC Cpg oligonucleotides can be useful in activating B cells, NK cells, and
CC antigen presenting cells, such as monocytes and macrophages. Cpg
CC oligonucleotides enhance antibody dependent cellular cytotoxicity and can
CC be used as an adjuvant in conjunction with tumour antigens to protect
CC against a tumour challenge.

SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match 100.0%; Score 20; DB 20; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 tccatgacgttcctgacgtt 20
|||||
Db 1 tccatgacgttcctgacgtt 20

```
RESULT 6
Z41949 ID Z41949 standard; DNA: 20 BP.
XX AC Z41949;
XX DT 24-JAN-2000 (first entry)
XX DE IL-12 secretion inducing CpG oligonucleotide 94.
XX DE
XX KW CpG oligonucleotide; phosphorothioate; interleukin-12; IL-12; secretion;
XX KW human PBMC; immune response; cancer; HIV; bacterial disease; asthma;
XX KW neoplastic disorder; Jaagsiekte; B cell; NK cell; ss; cytokine;
XX KW antigen presenting cell; infection; allergic disease.
XX OS Synthetic.
XX OS
XX PN WO9951259-A2.
XX PD 14-OCT-1999.
XX PD
XX PF 02-APR-1999; 99WO-US07335.
XX PR 03-APR-1998; 98US-0080729.
XX PR
XX PA (IOWA ) UNIV IOWA RES FOUND.
XX PI Kriegl AM, Welner G;
XX PI
XX DR WPI; 1999-620169/53.
XX DR
XX PT Novel synergistic combinations of immunostimulatory oligonucleotides
XX PT and immunopotentiating cytokines are useful for stimulating the immune
XX PT system -
XX PS Example 8; Page 89; 91pp; English.
XX PS
XX CC Sequences Z41856-Z41949 are phosphorothioate CpG oligonucleotides which
XX CC are used in the invention to induce interleukin-12 (IL-12) secretion
XX CC from human PBMC. The invention comprises stimulating an immune response
XX CC in a subject comprising administering to a subject exposed to an antigen,
XX CC an immunopotentiating cytokine and an immunostimulatory CpG
XX CC oligonucleotide to induce a synergistic antigen specific immune response.
XX CC The methods are useful for treating cancer by stimulating an antigen
XX CC specific immune response against a cancer antigen. The methods can also
XX CC be used to treat neoplastic disorders in humans, including but not
XX CC limited to: sarcoma, carcinoma, fibroma, lymphoma, melanoma,
XX CC neuroblastoma, retinoblastoma, and glioma. The methods are also useful
XX CC for treating infectious diseases, e.g. viral diseases such as HIV,
XX CC bacterial diseases, and fungal diseases. The methods may also be used to
XX CC treat allergic diseases, e.g. asthma. The methods and compositions may
XX CC also be applied to treat cancer and tumours in non human subjects,
XX CC e.g. cats and dogs. Neoplasias affecting agricultural livestock may also
XX CC be treated and include leukaemia, haemangioepithelioma and bovine ocular
XX CC neoplasia. Chronic, infectious, contagious diseases of sheep and goats
XX CC caused by the bacterium Corynebacterium pseudotuberculosis, and
XX CC contagious lung tumour of sheep caused by Jaagsiekte may also be treated.
XX CC CpG oligonucleotides can be useful in activating B cells, NK cells, and
XX CC antigen presenting cells, such as monocytes and macrophages. CpG
XX CC oligonucleotides enhance antibody dependent cellular cytotoxicity and can
XX CC be used as an adjuvant in conjunction with tumour antigens to protect
XX CC against a tumour challenge.
XX CC
XX SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;
```

```
Query Match 100.0%; Score 20; DB 20; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

QY 1 tccatgacgttcctgacgtt 20
 |||||
Db 1 tccatgacgttcctgacgtt 20
 ,

```
RESULT 7
Z28191 ID Z28191 standard; DNA: 20 BP.
XX AC Z28191;
XX DT 20-DEC-1999 (first entry)
XX DE Chlamydia trachomatis outer membrane protein gene-derived CpG oligo 4.
XX DE
XX KW Heart disease; inflammatory; autoimmune; cardiomyopathy; adjuvant;
XX KW CpG motif; vaccine; ds.
XX KW
XX OS Synthetic.
XX OS Chlamydia trachomatis.
XX OS
XX FH Key Location/Qualifiers
XX FT modified_base 1..20
XX FT /*tag= a
XX FT /mod_base= OTHER
XX FT /note= "OTHER = phosphorothioate linkage"
XX FT
XX PN US5962636-A.
XX PN
XX PD 05-OCT-1999.
XX PD
XX PF 12-AUG-1998; 98US-0133774.
XX PF
XX PR 12-AUG-1998; 98US-0133774.
XX PR
XX PA (AMGE-) AMGEN CANADA INC.
XX PA
XX PI Bachmaier K, Hessel AJ, Penninger JM, Neu N;
XX PI
XX DR WPI; 1999-589735/50.
XX DR
XX PT Peptides that induce or suppress inflammatory cardiomyopathy -
XX PT
XX PS Example 2; Column 25; 17pp; English.
XX PS
XX CC This sequence represents DNA encoding Chlamydia trachomatis 60 kD outer
XX CC membrane protein (OMP) gene-derived CpG oligonucleotide 4. This
XX CC oligonucleotide contains a CpG motif. It was tested for its ability to
XX CC act as an adjuvant for the M7A-alpha peptide (Y42723), which can induce
XX CC inflammatory cardiomyopathy (ICM) in mice. It was found to act as a
XX CC potent immunostimulator, whereas a oligonucleotide from the same
XX CC source which did not contain a CpG motif (Z28193) was hardly effective as
XX CC an adjuvant. Inflammatory cardiomyopathy peptides (Y42723, Y42725-Y42731)
XX CC can be used with such an adjuvant and an excipient in a vaccine for
XX CC decreasing ICM.
XX CC
XX SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;
```

```
Query Match 100.0%; Score 20; DB 20; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

QY 1 tccatgacgttcctgacgtt 20
 |||||
Db 1 tccatgacgttcctgacgtt 20

```
RESULT 8
X78802 ID X78802 standard; DNA: 20 BP.
XX AC X78802;
XX DT 06-SEP-1999 (first entry)
XX XX
```

DE HPV fusion protein Cpg oligonucleotide 1.
XX
KW Fusion protein; E6 protein; E7 protein; E6/E7; immunomodulator; tumour;
KW immunological fusion partner; Cpg oligonucleotide; immune response;
KW HPV antigen; prevention; treatment; primer; ss.
XX
OS Synthetic.
OS Human papillomavirus.
XX
PN WO9933868-A2.
XX
PD 08-JUL-1999.
XX
PF 18-DEC-1998; 98WO-EP08563.
XX
PR 24-DEC-1997; 97GB-0027262.
XX
PA (SMIK) SMITHKLINE BEECHAM BIOLOGICALS.
XX
PI Dalemans WLJ, Gerard CMG;
XX
DR WPI; 1999-405485/34.
XX
XX
PT Composition comprising an E6, E7 or E6/E7 fusion protein from HPV to
PT induce immune response to HPV
XX
PS Claim 11; Page 36; 62pp; English.
XX
CC X78791-X78801 represent nucleic acid sequences which encode novel
CC constructs comprising an E6 or E7 protein or E6/E7 fusion protein from
CC HPV (represented in Y25375-Y25386). These constructs are optionally
CC linked to an immunological fusion partner and an immunomodulatory Cpg
CC oligonucleotide. The products of the invention can be used to induce an
CC immune response in a patient to an HPV antigen. They can also be used
CC for preventing or treating HPV induced tumours. This sequence represents
CC a Cpg oligonucleotide which is used in the method of the invention.
XX
SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match 100.0%; Score 20; DB 20; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
1 |||||
Db 1 tccatgacgttcctgacgtt 20

RESULT 9
X88536
ID X88536 standard; DNA; 20 BP.
XX
AC X88536;
XX
DT 10-SEP-1999 (first entry)
XX
DE Cytosine-guanosine dinucleotide motif oligonucleotide #3.
XX
KW Cytosine-guanosine dinucleotide motif; Cpg; immunomodulation;
KW unethylated; vaccine; immunostimulation; immune response;
KW T-independent type 1 antigen; T-independent type 2 antigen;
KW polysaccharide conjugate antigen; ss.
XX
OS Synthetic.
XX
PN WO9933488-A2.
XX
PD 08-JUL-1999.
XX
PF 18-DEC-1998; 98WO-EP08562.
XX
PR 24-DEC-1997; 97GB-0027262.

XX
PA (SMIK) SMITHKLINE BEECHAM BIOLOGICALS.
XX
PI Dalemans WLJ, Laferriere CAJ, Prieeels J;
XX
DR WPI; 1999-405369/34.
XX
DR
XX
PT A vaccine composition for inducing a immune response to
PT T-independent type 1 or type 2 antigen or polysaccharide conjugate
PT antigen
XX
PS Claim 6; Page 31; 35pp; English.
XX
CC The present invention describes a formulation (A) comprising a
CC cytosine-guanosine dinucleotide motif (Cpg) oligonucleotide and
CC T-independent type 1 or type 2 antigens or polysaccharide conjugate
CC antigen. The present sequence represent a specifically claimed Cpg
CC oligonucleotide. A vaccine composition comprising the formulation is
CC used for inducing a immune response to T-independent type 1 or type 2
CC antigen or polysaccharide conjugate antigen. The use of
CC immunostimulatory Cpg oligonucleotide acts as an adjuvant to
CC pneumococcal polysaccharides.
XX
SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match 100.0%; Score 20; DB 20; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
1 |||||
Db 1 tccatgacgttcctgacgtt 20

RESULT 10
V74237
ID V74237 standard; DNA; 20 BP.
XX
AC V74237;
XX
DT 15-MAR-1999 (first entry)
XX
DE Cpg-N motif S-ODN 1826 DNA.
XX
KW Cpg-N motif; immunostimulation; antigen; Cpg-S motif; immunisation; ODN;
KW viral antigen; bacterial antigen; parasite; therapeutic; growth factor;
KW toxin; tumour suppressor; cytokine; apoptotic protein; interferon;
KW hormone; clotting factor; ligand; receptor; oligodeoxynucleotide; ss.
XX
OS Synthetic.
XX
PN WO9852581-A1.
XX
PD 26-NOV-1998.
XX
PF 20-MAY-1998; 98WO-US10408.
XX
PR 20-MAY-1997; 97US-0047233.
PR 20-MAY-1997; 97US-0047209.
XX
PA (OTTA-) OTTAWA CIVIC HOSPITAL, LOEB RES INST.
PA (QIAG-) QIAGEN GMBH.
PA (IOWA-) UNIV IOWA RES FOUND.
XX
PI Davis HL, Krieg AM, Schorr J, Wu T;
XX
DR WPI; 1999-059712/05.
XX
PT Use of neutralising Cpg and stimulating Cpg motifs in DNA vectors -
PT for enhancing the immunostimulatory effect of an antigen or
PT enhancing the expression of a therapeutic polypeptide


```
PS Example 1; Page 64; 109pp; English.
XX
CC V74237-V74253 are oligodeoxynucleotide (ODN) primers used to describe a
CC method for enhancing the immunostimulatory effect of an antigen encoded
CC by nucleic acid contained in a nucleic acid construct. The method
CC involves determining the Cpg-N and Cpg-S motifs present in the construct,
CC removing neutralising Cpg (Cpg-N) motifs and optionally inserting a
CC stimulatory Cpg (Cpg-S) motifs in the construct, thereby producing a
CC nucleic acid construct having enhanced immunostimulatory efficacy. The
CC method can be used for immunisation against viral antigens, e.g. from
CC hepatitis B virus (HBV), bacterial antigens or an antigen derived from a
CC parasite. They can also be used for expression of a therapeutic
CC polypeptide, e.g. growth factors, toxins, tumour suppressors, cytokines,
CC apoptotic proteins, interferons, hormones, clotting factors, ligands and
CC receptors.
XX
SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match          100.0%; Score 20; DB 20; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
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Db 1 tccatgacgttcctgacgtt 20

RESULT 11
V74242
ID V74242 standard; DNA; 20 BP.
AC V74242;
DT 15-MAR-1999 (first entry)
XX
DE Cpg-N motif O-ODN 2061 DNA.
XX
KW Cpg-N motif; immunostimulation; antigen; Cpg-S motif; immunisation; ODN;
KW viral antigen; bacterial antigen; parasite; therapeutic; growth factor;
KW toxin; tumour suppressor; cytokine; apoptotic protein; interferon;
KW hormone; clotting factor; ligand; receptor; oligodeoxynucleotide; ss.
XX
OS Synthetic.
XX
PN WO9852581-A1.
XX
PD 26-NOV-1998.
XX
PE 20-MAY-1998; 98WO-US10408.
XX
PR 20-MAY-1997; 97US-0047233.
PR 20-MAY-1997; 97US-0047209.
XX
PA (OTTA-) OTTAWA CIVIC HOSPITAL LOEB RES INST.
PA (QIAG-) QIAGEN GMBH.
PA (IOWA-) UNIV IOWA RES FOUND.
XX
PI Davis HL, Krieg AM, Schorr J, Wu T;
XX
DR WPI; 1999-059712/05.
XX
PT Use of neutralising Cpg and stimulating Cpg motifs in DNA vectors -
PT for enhancing the immunostimulatory effect of an antigen or
PT enhancing the expression of a therapeutic polypeptide
XX
PS Example 1; Page 64; 109pp; English.
XX
CC V74237-V74253 are oligodeoxynucleotide (ODN) primers used to describe a
CC method for enhancing the immunostimulatory effect of an antigen encoded
CC by nucleic acid contained in a nucleic acid construct. The method
CC involves determining the Cpg-N and Cpg-S motifs present in the construct,
CC removing neutralising Cpg (Cpg-N) motifs and optionally inserting a
CC stimulatory Cpg (Cpg-S) motifs in the construct, thereby producing a
CC nucleic acid construct having enhanced immunostimulatory efficacy. The
CC method can be used for immunisation against viral antigens, e.g. from
CC hepatitis B virus (HBV), bacterial antigens or an antigen derived from a
CC parasite. They can also be used for expression of a therapeutic
CC polypeptide, e.g. growth factors, toxins, tumour suppressors, cytokines,
CC apoptotic proteins, interferons, hormones, clotting factors, ligands and
CC removing neutralising Cpg (Cpg-N) motifs and optionally inserting
```

```
CC stimulatory Cpg (Cpg-S) motifs in the construct, thereby producing a
CC nucleic acid construct having enhanced immunostimulatory efficacy. The
CC method can be used for immunisation against viral antigens, e.g. from
CC hepatitis B virus (HBV), bacterial antigens or an antigen derived from a
CC parasite. They can also be used for expression of a therapeutic
CC polypeptide, e.g. growth factors, toxins, tumour suppressors, cytokines,
CC apoptotic proteins, interferons, hormones, clotting factors, ligands and
CC receptors.
XX
SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match          100.0%; Score 20; DB 20; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
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Db 1 tccatgacgttcctgacgtt 20

RESULT 12
V74244
ID V74244 standard; DNA; 20 BP.
XX
AC V74244;
XX
DT 15-MAR-1999 (first entry)
XX
DE Cpg-N motif SOS-ODN 1980 DNA.
XX
KW Cpg-N motif; immunostimulation; antigen; Cpg-S motif; immunisation; ODN;
KW viral antigen; bacterial antigen; parasite; therapeutic; growth factor;
KW toxin; tumour suppressor; cytokine; apoptotic protein; interferon;
KW hormone; clotting factor; ligand; receptor; oligodeoxynucleotide; ss.
XX
OS Synthetic.
XX
PN WO9852581-A1.
XX
PD 26-NOV-1998.
XX
PE 20-MAY-1998; 98WO-US10408.
XX
PR 20-MAY-1997; 97US-0047233.
PR 20-MAY-1997; 97US-0047209.
XX
PA (OTTA-) OTTAWA CIVIC HOSPITAL LOEB RES INST.
PA (QIAG-) QIAGEN GMBH.
PA (IOWA-) UNIV IOWA RES FOUND.
XX
PI Davis HL, Krieg AM, Schorr J, Wu T;
XX
DR WPI; 1999-059712/05.
XX
PT Use of neutralising Cpg and stimulating Cpg motifs in DNA vectors -
PT for enhancing the immunostimulatory effect of an antigen or
PT enhancing the expression of a therapeutic polypeptide
XX
PS Example 1; Page 64; 109pp; English.
XX
CC V74237-V74253 are oligodeoxynucleotide (ODN) primers used to describe a
CC method for enhancing the immunostimulatory effect of an antigen encoded
CC by nucleic acid contained in a nucleic acid construct. The method
CC involves determining the Cpg-N and Cpg-S motifs present in the construct,
CC removing neutralising Cpg (Cpg-N) motifs and optionally inserting a
CC stimulatory Cpg (Cpg-S) motifs in the construct, thereby producing a
CC nucleic acid construct having enhanced immunostimulatory efficacy. The
CC method can be used for immunisation against viral antigens, e.g. from
CC hepatitis B virus (HBV), bacterial antigens or an antigen derived from a
CC parasite. They can also be used for expression of a therapeutic
CC polypeptide, e.g. growth factors, toxins, tumour suppressors, cytokines,
CC apoptotic proteins, interferons, hormones, clotting factors, ligands and
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CC receptors.
XX
SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match
Best Local Similarity 100.0%; Score 20; DB 20; Length 20;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 1 tccatgacgttcctgacgtt 20

RESULT 13
Z99004
ID Z99004 standard; DNA; 20 BP.
XX
AC Z99004;
XX
XX 21-JUN-2000 (first entry)
XX
DE Cpg motif for immunostimulatory oligonucleotide 1826.
XX
XX
KW Immunoprotective; vaccine; antigen; saponin adjuvant; immune response;
KW Immunostimulatory oligonucleotide; unmethylated Cpg dinucleotide;
KW mammal; human; animal; ss.
XX
XX
OS Synthetic.
XX
PN WO200009159-A1.
XX
PD 24-FEB-2000.
XX
PF 06-AUG-1999; 99WO-US17956.
XX
PR 10-AUG-1998; 98US-0095913.
PR 08-APR-1999; 99US-0128608.
XX
PA (AQUIT-) AQUILA BIOPHARMACEUTICALS INC.
XX
PI Kensil CA;
XX
DR WPI; 2000-224181/19.
XX
PT A vaccine composition comprising an antigen, saponin adjuvant and
PT immunostimulatory Cpg oligonucleotide, useful for stimulating immunity
PT and increasing immune responses -
XX
PS Claim 10; Page 19; 38pp; English.
XX
XX The invention relates to a vaccine composition comprising an antigen,
CC a saponin adjuvant and an immunostimulatory oligonucleotide. The
CC immunostimulatory oligonucleotide preferably comprises at least one
CC unmethylated Cpg dinucleotide. This sequence represents an example of
CC the immunostimulatory oligonucleotide. The vaccine composition increases
CC the immune response to the antigen when administered to a mammal,
CC especially a human or animal. It further stimulates immunity and
CC especially enhances antibody production to the antigen, preferably in a
CC positive synergistic manner. It further enhances cell-mediated immunity.
CC The immune adjuvant in particular can be used to increase the immune
CC response to an antigen in an individual or a test system.
XX
SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match
Best Local Similarity 100.0%; Score 20; DB 21; Length 20;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 tccatgacgttcctgacgtt 20
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Db 1 tccatgacgttcctgacgtt 20

RESULT 14
Z99174
ID Z99174 standard; DNA; 20 BP.
XX
AC Z99174;
XX
XX 21-JUN-2000 (first entry)
XX
DE Inflammatory cardiomyopathy immunostimulatory oligonucleotide #3.
XX
KW Cardiant; murine alpha myosin heavy chain; inflammatory myocarditis;
KW autoimmune inflammatory cardiomyopathy; Chlamydia; antibody; vaccine;
KW hybridization probe; immunostimulatory; ss.
XX
XX
OS Synthetic.
XX
PN US6034230-A.
XX
PD 07-MAR-2000.
XX
PF 03-MAY-1999; 99US-0303862.
XX
PR 12-AUG-1998; 98US-0133774.
XX
PA (AMGE-) AMGEN CANADA INC.
XX
PI Neu N, Penninger JM, Bachmaier K, Hessel AJ;
XX
DR WPI; 2000-255712/22.
XX
PT DNA molecules encoding novel myocardial peptides used for inhibiting
PT and inducing inflammatory cardiomyopathy in vivo -
XX
PS Disclosure; Column 17; 17pp; English.
XX
XX The invention relates to the isolation of sequences coding for peptide
CC sequences derived from bacteria and viruses which may cause inflammatory
CC cardiomyopathy. The peptide sequences are searched based on the sequence
CC of the M7A peptides derived from the murine alpha myosin heavy chain
CC polypeptide. The consensus sequence of the murine M7A-alpha/beta peptides
CC (Y83813) was used to search the PIR public database for similar bacterial
CC and viral sequences able to cause inflammatory cardiomyopathy. The screen
CC isolated the peptides Y83814-Y83819 and their corresponding coding
CC sequences Z99164-Z99169. The peptides encoded by the DNAs are used, alone
CC or in conjunction with other therapeutics, for inducing or inhibiting
CC inflammatory cardiomyopathy in vivo, where the cardiomyopathy is
CC autoimmune inflammatory cardiomyopathy, and inflammatory cardiomyopathy
CC caused by Chlamydia or other bacterial or viral infections that cause
CC inflammatory cardiomyopathy. The oligonucleotides Z99172-Z99176 were
CC shown to increase the immunogenicity of the immunostimulatory peptides
CC when injected simultaneously. The peptides may also be used for
CC increasing inflammatory myocarditis in a mammal. Antibodies against the
CC peptides and the peptides themselves are used for measuring the risk of
CC inflammatory cardiomyopathy in a mammal. The peptides may also be used
CC in vaccines. Nucleic acids encoding the peptides may be used as
CC hybridization probes, e.g. in diagnostic assays to test for the
CC presence of Chlamydia DNA.
XX
SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match
Best Local Similarity 100.0%; Score 20; DB 21; Length 20;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 tccatgacgttcctgacgtt 20
 |||||
Db 1 tccatgacgttcctgacgtt 20

RESULT 15

261010
ID 261010 standard; DNA; 20 BP.
XX
AC 261010;
XX
DT 30-MAY-2000 (first entry)
XX
DE Nucleotide sequence of an immunostimulatory Cpg oligonucleotide.
XX
KW Immunostimulatory; stereoisomer; Cpg oligonucleotide; Th2; Th1; asthma;
KW allergic reaction; allergen; cancer antigen; cancer; immunoinhibitory;
KW inflammatory disease; inflammatory bowel disease; autoimmune disease;
KW gingivitis; psoriasis; sepsis; ss.
XX
OS Synthetic.
XX
PN WO200006588-A1.
XX
PD 10-FEB-2000.
XX
PE 27-JUL-1999; 99WO-US17100.
XX
PR 27-JUL-1998; 98US-0094370.
XX
PA (IOWA) UNIV IOWA RES FOUND.
PA (CPGI-) CPG IMMUNOPHARMACEUTICALS INC.
XX
PI Kriegl AM;
XX
DR WPI; 2000-195254/17.
XX
PT Immunostimulatory and immunoinhibitory stereoisomers of Cpg
PT oligonucleotides useful for immunotherapy of cancer -
XX
PS Disclosure; Page 12; 88pp; English.
XX
XX 260933-261015 represent immunostimulatory stereoisomers of Cpg
CC oligonucleotides. The sequences are derived from generic nucleic
CC acid sequence, from which immunoinhibitory sequences may also be
CC derived. The immunostimulatory nucleic acids can be co-administered
CC with an antigen to induce an antigen-specific immune response. The
CC immunostimulatory nucleic acids can also be used in methods for
CC redirecting a subject's immune response from a Th2 to a Th1, for
CC treating asthma, for desensitizing a subject against the occurrence
CC of an allergic reaction in response to contact with an allergen, for
CC activating an immune cell, especially a lymphocyte or a dendritic cell
CC expressing a cancer antigen or for treating cancer. The immunoinhibitory
CC nucleic acid can be used to prevent an immune response, especially where
CC the immune response in the subject is excessive due to having received
CC an immune stimulating compound. The immunoinhibitory nucleic acid can
CC be used to treat a subject having or at risk of an inflammatory disease,
CC especially inflammatory bowel disease, autoimmune disease, gingivitis,
CC psoriasis and sepsis.
XX
SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;
QY 1 tccatgacgttcctgacgtt 20
Db 1 tccatgacgttcctgacgtt 20
Query Match 100.0%; Score 20; DB 21; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Search completed: December 4, 2000, 21:09:55
Job time: 16673 sec

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DEFINITION      Sequence 8 from patent US 5846757.
ACCESSION       AR063883
VERSION         AR063883.1  GI:5993191
KEYWORDS
SOURCE          Unknown.
ORGANISM        Unknown.
                Unclassified.
REFERENCE       1 (bases 1 to 7175)
AUTHORS        Harpold,M.M., Ellis,S.B., Williams,M.E., Feldman,D.H., McCue,A.F.
                and Brenner,R.
TITLE          Human calcium channel .alpha..sub.1, .alpha. .sub.2, and .beta.
                subunits and assays using them
JOURNAL         Patent: US 5846757-A 8 08-DEC-1998;
FEATURES       Location/Qualifiers
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                /organism="unknown"
BASE COUNT     1415 a 2197 c 2168 g 1395 t
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Query Match      84.0%; Score 16.8; DB 5; Length 7175;
Best Local Similarity 90.0%; Pred. No. 2.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
    |||||
Db 652 TCCATGACGTTCCAGCCGTT 633

RESULT 15
AR067883/c
LOCUS           AR067883      7175 bp      DNA           PAT      29-SEP-1999
DEFINITION     Sequence 8 from patent US 5851824.
ACCESSION      AR067883
VERSION        AR067883.1  GI:5999105
KEYWORDS
SOURCE         Unknown.
ORGANISM       Unknown.
                Unclassified.
REFERENCE      1 (bases 1 to 7175)
AUTHORS        Harpold,M.M., Ellis,S.B., Williams,M.E., Feldman,D.H., McCue,A.F.
                and Brenner,R.
TITLE          Human calcium channel .alpha.-1C/.alpha.-1D, .alpha.-2, .beta.-1,
                and .gamma.subunits and cells expressing the DNA
JOURNAL         Patent: US 5851824-A 8 22-DEC-1998;
FEATURES       Location/Qualifiers
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                /organism="unknown"
BASE COUNT     1415 a 2197 c 2168 g 1395 t
ORIGIN

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Best Local Similarity 90.0%; Pred. No. 2.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
    |||||
Db 652 TCCATGACGTTCCAGCCGTT 633

Search completed: December 4, 2000, 20:47:28
Job time: 18088 sec
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Db	652	TCCATGACGTTCCAGCGTT	633		
RESULT	12				
D86600/c					
LOCUS	D86600	7113 bp	mrna	INV	08-MAR-2000
DEFINITION	Loligo bleekeri mRNA for voltage-dependent calcium channel, complete cds.				
ACCESSION	D86600				
VERSION	D86600.2	GI:7209875			
KEYWORDS	voltage-dependent calcium channel; neural calcium channel alphanal-subunit.				
SOURCE	Loligo bleekeri cDNA to mRNA.				
ORGANISM	Loligo bleekeri				
	Eukaryota; Metazoa; Mollusca; Cephalopoda; Coleoidea; Teuthoida; Myopsida; Loliginidae; Loligo.				
REFERENCE	1 (sites)				
AUTHORS	Kimura,T., Shouno,O., Hirota,K., Saito,T., Matsumoto,G. and Sato,C.				
TITLE	Molecular cloning and characterization of a putative neural calcium channel alphanal-subunit from squid optic lobe				
JOURNAL	Biochem. Biophys. Res. Commun. 230 (1), 147-154 (1997)				
MEDLINE	97148591				
REFERENCE	2 (bases 1 to 7113)				
AUTHORS	Kimura,T., Shouno,O., Sato,C., Hirota,K., Hanyu,Y., Saito,T. and Matsumoto,G.				
TITLE	Primary structure of a putative calcium channel alphanal-subunit from squid optic lobe				
JOURNAL	Unpublished (1996)				
REFERENCE	3 (bases 1 to 7113)				
AUTHORS	Kimura,T.				
TITLE	Direct Submission				
JOURNAL	Submitted (22-JUL-1996) to the DDBJ/EMBL/GenBank databases. Tadashi Kimura, Electrotechnical Laboratory, Supermolecular Science Division; 1-1-4 Umezono, Tsukuba, Ibaraki 305, Japan (E-mail:eveda@etl.go.jp, Tel:029854508066616)				
COMMENT	On Mar 8, 2000 this sequence version replaced gi:1817549. Sequence updated (27-Nov-1996) by: Tadashi Kimura				
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	/db_xref="taxon:6617"				
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	/protein_id="BAA13136.2"				
	/db_xref="GI:7209876"				
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	VLSGEFAKERERVENRAFLKLRQQQIERLNGYLEWICKAAEEVILDEERKKDDGTI				
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	LKNRKAHGRCAGFWRAEKHLRFTIRKCVKTQGFYWFVILVFLNLCVASEHYGQAEW				
	HTEFLYVMEFAFLALFMSEMLIKMYGLGVRVLVQSSFNFDVCVILVSIIEVIWSAIK				
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NSEEPAFGPKPMLPYSSMFIPTNPVRRFCHFVFDLFIMIVICASSIALAA
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ICALVAFAGDAAGNLNTIKSLRVLRLPKTIINRIPKLKAFLDCVAVNSLKNVSNV
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INRHNNERSFFYAVLLFCATGESWQOIMLSCLSGRPCDPESKMLDNSCGLDIAYIY
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EDDDWC"

BASE COUNT 2146 a 1383 c 1514 g 2070 t
ORIGIN

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Best Local Similarity 90.0%; Pred. No. 2.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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||||| ||||| |||||
Db 747 TCCATGATGTTCCAGACGTT 728

RESULT 13
AR022380/c
LOCUS AR022380 7175 bp DNA PAT 05-DEC-1998
DEFINITION Sequence 8 from patent US 5792846.
ACCESSION AR022380
VERSION AR022380.1 GI:3976442

KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
FEATURES
source
BASE COUNT 1415 a 2197 c 2168 g 1395 t
ORIGIN

Query Match 84.0%; Score 16.8; DB 5; Length 7175;
Best Local Similarity 90.0%; Pred. No. 2.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
||||| ||||| |||||
Db 652 TCCATGACGTTCCAGCGTT 633

RESULT 14
AR063883/c
LOCUS AR063883 7175 bp DNA PAT 29-SEP-1999

RESULT 7
A90870 LOCUS A90870 20 bp DNA PAT 22-JAN-2000
DEFINITION Sequence 5 from Patent EP0855184.
ACCESSION A90870
VERSION A90870.1 GI:6739264
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 20)
AUTHORS Heeg,K.P. and Lipford,G.B.
TITLE Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination
JOURNAL Patent: EP 0855184-A 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
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Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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Db 1 TCCATGACGTTCCCTGATGCT 20

RESULT 8
A93512 LOCUS A93512 20 bp DNA PAT 22-JAN-2000
DEFINITION Sequence 5 from Patent WO9740163.
ACCESSION A93512
VERSION A93512.1 GI:6741731
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 20)
AUTHORS Colpan,M. and Schorr,J.
TITLE NUCLEIC ACID VACCINATION FOR PARVOVIRAL INFECTIONS
JOURNAL Patent: WO 9740163-A 30-OCT-1997;
COLPAN METIN (DE); SCHORR JOACHIM (DE)
FEATURES
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/db_xref="taxon:32644" 7 t
BASE COUNT 3 a 6 c 4 g 7 t
ORIGIN

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Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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|||||
Db 1 TCCATGACGTTCCCTGATGCT 20

RESULT 9
A93521 LOCUS A93521 20 bp DNA PAT 22-JAN-2000
DEFINITION Sequence 14 from Patent WO9740163.
ACCESSION A93521
VERSION A93521.1 GI:6741738
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 20)
AUTHORS Colpan,M. and Schorr,J.
TITLE NUCLEIC ACID VACCINATION FOR PARVOVIRAL INFECTIONS
JOURNAL Patent: WO 9740163-A 30-OCT-1997;
COLPAN METIN (DE); SCHORR JOACHIM (DE)
FEATURES
source Location/Qualifiers
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Best Local Similarity 90.0%; Pred. NO. 2.2e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
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Db 1 TCCATGACGTTCCCTGATGCT 20

ORGANISM unidentified
unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Colpan,M. and Schorr,J.
TITLE NUCLEIC ACID VACCINATION FOR PARVOVIRAL INFECTIONS
JOURNAL Patent: WO 9740163-A 30-OCT-1997;
COLPAN METIN (DE); SCHORR JOACHIM (DE)
FEATURES
source Location/Qualifiers
1. .20
/organism="unidentified"
/db_xref="taxon:32644" 7 t
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ORIGIN

Query Match 84.0%; Score 16.8; DB 5; Length 20;
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Db 1 TCCATGACGTTCCCTGATGCT 20

RESULT 10
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DEFINITION Sequence 14 from patent US 5429921.
ACCESSION I12881
VERSION I12881.1 GI:910858
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 2470)
AUTHORS Harpold,M.M., Ellis,S.B., Williams,M.E., Feldman,D.H., McCue,A.F. and Brenner,R.
TITLE Assays for agonists and antagonists of recombinant human calcium channels
JOURNAL Patent: US 5429921-A 14 04-JUL-1995;
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source Location/Qualifiers
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RESULT 11
I12880/c LOCUS I12880 5467 bp DNA PAT 26-JUL-1995
DEFINITION Sequence 12 from patent US 5429921.
ACCESSION I12880
VERSION I12880.1 GI:910857
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 5467)
AUTHORS Harpold,M.M., Ellis,S.B., Williams,M.E., Feldman,D.H., McCue,A.F. and Brenner,R.
TITLE Assays for agonists and antagonists of recombinant human calcium channels
JOURNAL Patent: US 5429921-A 12 04-JUL-1995;
FEATURES
source Location/Qualifiers

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RESULT 3
AB016662S3/c
LOCUS AB016662S3 595 bp DNA ROD 14-APR-2000
DEFINITION Mus musculus gene for aldose reductase, exon 9.
ACCESSION AB016664
VERSION AB016664.1 GI:4586467
KEYWORDS aldose reductase.
SEGMENT 3 of 4
SOURCE Mus musculus (strain:BALB/c) DNA.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (sites)
AUTHORS Li,H., Nobukuni,Y., Gui,T. and Yabe-Nishimura,C.
TITLE Characterization of genomic regions directing the cell-specific
expression of the mouse aldose reductase gene
JOURNAL Biochem. Biophys. Res. Commun. 255 (3), 759-764 (1999)
MEDLINE 99160426
REFERENCE 2 (bases 1 to 595)
AUTHORS Yabe-Nishimura,C. and Li,H.
TITLE Direct Submission
JOURNAL Submitted (31-JUL-1998) to the DDBJ/EMBL/GenBank databases. Chihiro
Yabe-Nishimura, Kyoto Prefectural University of Medicine,
Department of Pharmacology; Kawaramachi-Hirokoji, Kamikyoku, Kyoto,
Kyoto 602-8566, Japan (E-mail:nchihiro@basic.kpu-m.ac.jp,
Tel:+81-75-251-5333, Fax:+81-75-251-5348)
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RESULT 4
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ACCESSION A89782
VERSION A89782.1 GI:6738296
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Lipford,G.B. and Heeg,K.
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
JOURNAL Patent: WO 9832462-A 30-JUL-1998;
LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
FEATURES
source Location/Qualifiers
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Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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Db 1 TCCATGACGTTCTGATGCT 20

RESULT 5
A89783
LOCUS A89783 20 bp DNA PAT 22-JAN-2000
DEFINITION Sequence 5 from Patent WO9832462.
ACCESSION A89783
VERSION A89783.1 GI:6738297
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Lipford,G.B. and Heeg,K.
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
JOURNAL Patent: WO 9832462-A 30-JUL-1998;
LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
FEATURES
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RESULT 6
A90869
LOCUS A90869 20 bp DNA PAT 22-JAN-2000
DEFINITION Sequence 4 from Patent EP0855184.
ACCESSION A90869
VERSION A90869.1 GI:6739263
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Heeg,K.P. and Lipford,G.B.
TITLE Pharmaceutical composition comprising a polynucleotide and an
antigen especially for vaccination
JOURNAL Patent: EP 0855184-A 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
FEATURES
source Location/Qualifiers
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Best Local Similarity 90.0%; Pred. No. 2.2e+02;
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Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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Db 1 TCCATGACGTTCTGATGCT 20

RESULT 5
A89783
LOCUS A89783 20 bp DNA PAT 22-JAN-2000
DEFINITION Sequence 5 from Patent WO9832462.
ACCESSION A89783
VERSION A89783.1 GI:6738297
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Lipford,G.B. and Heeg,K.
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
JOURNAL Patent: WO 9832462-A 30-JUL-1998;
LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
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Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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Db 1 TCCATGACGTTCTGATGCT 20

RESULT 6
A90869
LOCUS A90869 20 bp DNA PAT 22-JAN-2000
DEFINITION Sequence 4 from Patent EP0855184.
ACCESSION A90869
VERSION A90869.1 GI:6739263
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Heeg,K.P. and Lipford,G.B.
TITLE Pharmaceutical composition comprising a polynucleotide and an
antigen especially for vaccination
JOURNAL Patent: EP 0855184-A 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
FEATURES
source Location/Qualifiers
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BASE COUNT 3 a 6 c 4 g 7 t
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Best Local Similarity 90.0%; Pred. No. 2.2e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 tccatgacgttcctgacgtt 20
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C 17	16.8	84.0	7362	5	AR022379	Sequence
C 18	16.8	84.0	7362	5	AR063882	Sequence
C 19	16.8	84.0	7362	5	AR067882	Sequence
C 20	16.8	84.0	7364	67	HUMCACHNT	
C 21	16.8	84.0	7713	3	RABCCBIII	
C 22	16.8	84.0	30713	32	CELW05G11	
C 23	16.8	84.0	69051	83	AC073171	Homo sapi
C 24	16.8	84.0	124595	90	HSJ322E17	
C 25	16.8	84.0	148871	78	AC025324	Homo sapi
C 26	16.8	84.0	170595	78	AC025937	Homo sapi
C 27	16.8	84.0	186526	69	AC008669	Homo sapi
C 28	16.4	82.0	3694	12	AF224508	
C 29	16	80.0	2271	7	AF135189	Mus muscu
C 30	16	80.0	82686	7	AC004669	Arabidops
C 31	16	80.0	133137	10	AC007245	
C 32	15.8	79.0	371	82	AC062158	Homo sapi
C 33	15.8	79.0	615	81	AC048760	Giardia i
C 34	15.8	79.0	720	8	CNS01CTC	
C 35	15.8	79.0	758	85	AC076687	Botrytis
C 36	15.8	79.0	762	81	AC051209	Giardia i
C 37	15.8	79.0	768	81	AC055639	Giardia i
C 38	15.8	79.0	777	84	AC075866	Giardia i
C 39	15.8	79.0	790	80	AC037305	Giardia i
C 40	15.8	79.0	799	81	AC058159	Giardia i
C 41	15.8	79.0	823	80	AC039484	Giardia i
C 42	15.8	79.0	873	81	AC049019	Giardia i
C 43	15.8	79.0	1084	81	AC048024	Giardia i
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ALIGNMENTS

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LOCUS	Expression vector pMCG16-S,					complete sequence.
DEFINITION	AF053408					
ACCESSION	AF053408.1	GI:3746485				
VERSION						
KEYWORDS						
SOURCE	Expression vector pMCG16-S.					
ORGANISM	Expression vector pMCG16-S.					
REFERENCE	1 (bases 1 to 4227)					
AUTHORS	Krieg,A.M., Wu,T., Weeratna,R., Efler,S.M., Love-Homan,L., Yang,L.,					
TITLE	Sequence motifs in adenoviral DNA block immune activation by					
JOURNAL	stimulatory CpG motifs					
MEDLINE	Proc. Natl. Acad. Sci. U.S.A. 95 (21), 12631-12636 (1998)					
REFERENCE	2 (bases 1 to 4227)					
AUTHORS	Wu,T., Efler,S.M., Davis,H.L., Krieg,A.M. and Schorr,J.					
TITLE	Direct Submission					
JOURNAL	Submitted (12-MAR-1998) HGD, Loeb Health Research Institute, 725					
FEATURES	Parkdale Ave., Ottawa, ON K1Y 4E9, Canada					
source	Location/Qualifiers					
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DEFINITION	Expression vector pMCG50-S,				17-OCT-1998
ACCESSION	AF053409				
VERSION	AF053409.1	GI:3746487			
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SOURCE	Expression vector pMCG50-S.				
ORGANISM	Expression vector pMCG50-S.				
REFERENCE	1 (bases 1 to 4625)				
AUTHORS	Krieg,A.M., Wu,T., Weeratna,R., Efler,S.M., Love-Homan,L., Yang,L.,				
TITLE	Sequence motifs in adenoviral DNA block immune activation by				
JOURNAL	stimulatory CpG motifs				
MEDLINE	Proc. Natl. Acad. Sci. U.S.A. 95 (21), 12631-12636 (1998)				
REFERENCE	2 (bases 1 to 4625)				
AUTHORS	Wu,T., Efler,S.M., Davis,H.L., Krieg,A.M. and Schorr,J.				
TITLE	Direct Submission				
JOURNAL	Submitted (12-MAR-1998) HGD, Loeb Health Research Institute, 725				
FEATURES	Parkdale Ave., Ottawa, ON K1Y 4E9, Canada				
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QY	1	tccatgacgttcctgacgtt	20		
Db	2444	TCCATGACGTTCTTGACGTT	2463		

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

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87.962 Million cell updates/sec

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Perfect score: 20
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Scoring table: IDENTITY_NUC
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Searched: 1033670 seqs, 2183789903 residues
Total number of hits satisfying chosen parameters: 2067340

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Post-processing: Minimum Match 0%
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- 92: gb_sts2:*
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- 94: gb_vl2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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	4	16.8	84.0	20 5	A89782	A89782 Sequence 4
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7	16.8	84.0	20 5	A90870	A90870 Sequence 5	
8	16.8	84.0	20 5	A93512	A93512 Sequence 5	
9	16.8	84.0	20 5	A93521	A93521 Sequence 14	
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